

STIC Search Report Biotech-Chem Library

STIC Dalabase Tracking Number 1988

TO: Ralph J Gitomer

Location: rem/3D65/3C18

Art Unit: 1655

Thursday, November 03, 2005

Case Serial Number: 10/750637

From: Alex Waclawiw

Location: Biotech-Chem Library

Rem 1A71

Phone: 272-2534

Alexandra.waclawiw@uspto.gov

Search Notes			
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	<u>,</u>		



Access DB# 169613

SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: 261	TOMER	Examiner # :	69630 D	ate: 16/26	105	-
Art Unit: 655 Phone N	lumber 30	Serial Nu	mber: /0/75	0,637	•	_
Mail Box and Bldg/Room Location	:	Results Format Prefe	erred (circle): P	APER DIS	K E-N	4AIL
305/3C18				1		M9
If more than one search is subm	iπea, piease.pri ************	OFITIZE SEAFCRES IN (order or need	l. *******	*****	****
Please provide a detailed statement of the Include the elected species or structures, k utility of the invention. Define any terms known. Please attach a copy of the cover s	eywords, synonyms, that may have a spec	acronyms, and registry rail meaning. Give exam	numbers, and com	bine with the o	concept	or
Title of Invention:	·		~	,		
Inventors (please provide full names):	•					
Earliest Priority Filing Date:						
For Sequence Searches Only Please include appropriate serial number.	le all pertinent inform	ation (parent, child, divisio	nal, or issued pate	nt numbers) alo	ng with I	the
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STAFF USE ONLY	Type of Search	· Vendo	rs and cost wher	e applicable		
Searcher:	'NA Sequence (#)	STN	\$	•		
Searcher Phone #: Point of Contact: Alexandra Waclawiw	AA Sequence (#)	Dialog	-/ .			
Searcher Location choical Info. Specialist	Structure (#)	-Questel/Orbit				
Date Searcher Picked Up: 1/2	· Bibliographic	Dr.Link		· · · · · · · · · · · · · · · · · · ·		
Date Completed: 11-3	Litigation	Lexis/Nexis				
"Searcher Prep'& Review Time:	Fulltext	'Sequence Systems	*		· · ·	•
Clerical Prep Time:	Patent Family	-WWW/Internet				
Online Time: 7.0	. Other	Other (specify)				

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(FILE 'REGISTRY' ENTERED AT 13:43:55 ON 03 NOV 2005)
               DEL HIS Y
               D SAVE
               ACT ENZYMES/A
              1) SEA ABB=ON PLU=ON
                                   "ORGANOPHOSPHORUS HYDROLASE"/CN
L1
              1) SEA ABB=ON PLU=ON
                                    "ORGANOPHOSPHORUS ACID ANHYDROLASE"/CN
L2
              1) SEA ABB=ON PLU=ON DFPASE/CN
L3
              1) SEA ABB=ON PLU=ON PHOSPHOTRIESTERASE/CN
L4
              3 SEA ABB=ON PLU=ON (L1 OR L2 OR L3 OR L4)
L5
               ACT POLYELECTRO/A
              4 SEA ABB=ON PLU=ON 9002-98-6 OR 9003-01-4 OR 49718-51-6 OR
L6
                26062-79-3 OR 26062-79-3
               ACT ENDCAP/A
               _____
              3 SEA ABB=ON PLU=ON 5919-74-4 OR 149305-62-4 OR 1760-24-3
L7
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           1505 SEA ABB=ON PLU=ON L5
L8
L9
           6210 SEA ABB=ON
                           PLU=ON
                                   ENZYMES, USES/CT
                           PLU=ON
L10
          31031 SEA ABB=ON
                                   L6
                                   POLYELECTROLYTES/CT
                           PLU=ON
L11
          18108 SEA ABB=ON
                           PLU=ON
                                   END CAP?/OBI
L12
           1903 SEA ABB=ON
           4507 SEA ABB=ON
                           PLU=ON
                                   L7
L13
           7698 SEA ABB=ON
                           PLU=ON
                                   L8 OR L9
L14
                                   L10 OR L11
L15
          46227 SEA ABB=ON
                           PLU=ON
                                  L12 OR L13
           6408 SEA ABB=ON
                           PLU=ON
L16
                                   L14 AND L15 AND L16
                           PLU=ON
L17
              2 SEA ABB=ON
                                   L14 AND L15
L18
            146 SEA ABB=ON
                           PLU=ON
                                    TOXIN#/OBI OR CHEMICAL#/OBI
        1188205 SEA ABB=ON
                           PLU=ON
L19
L20
             16 SEA ABB=ON PLU=ON
                                    L18 AND L19
                                    MATERIAL/OBI OR BIOACTIV?/OBI OR GLASS/OBI
L21
        2314372 SEA ABB=ON PLU=ON
                OR SILICA/OBI OR TEXTILE#/OBI OR CLOTH?/OBI OR THREAD?/OBI
L22
             66 SEA ABB=ON PLU=ON
                                   L18 AND L21
                                   DECONTAMI?/OBI
L23
          10530 SEA ABB=ON
                           PLU=ON
L24
         737361 SEA ABB=ON PLU=ON
                                   CATALYS?/OBI
              3 SEA ABB=ON PLU=ON L22 AND L23
L25
                           PLU=ON L22 AND L24
L26
             20 SEA ABB=ON
                           PLU=ON
                                   L17 OR L25
L27
              4 SEA ABB=ON
                                   L26 NOT L27
L28
             19 SEA ABB=ON
                           PLU=ON
           7384 SEA ABB=ON PLU=ON
                                   ENZYME/OBI (L) MODIF?/OBI
L29
                                   L21 (L) L29
L30
            160 SEA ABB=ON
                           PLU=ON
                                   L30 AND L19
                           PLU=ON
L31
             17 SEA ABB=ON
                           PLU=ON PROTECT?/OBI OR DECONTAMIN?/OBI OR
L32
         950582 SEA ABB=ON
                SAFETY/OBI OR COAT?/OBI
                           PLU=ON L31 AND L32
L33
              4 SEA ABB=ON
                            PLU=ON CHELAT?/OBI (L) (METAL/OBI OR CU/OBI OR
L34
          21409 SEA ABB=ON
                COPPER/OBI)
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                E COPPER/CN
L35
              1 SEA ABB=ON PLU=ON COPPER/CN
     FILE 'CAPLUS' ENTERED AT 13:58:51 ON 03 NOV 2005
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L36	494865	SEA ABB=ON	PLU=ON	L35
L37	3	SEA ABB=ON	PLU=ON	L30 AND (L36 OR L34)
L38	9	SEA ABB=ON	PLU=ON	L18 AND (L34 OR L36)
L39	17	SEA ABB=ON	PLU=ON	L37 OR L38 OR L33 OR L27
L40	64633	SEA ABB=ON	PLU=ON	TOXINS/CT
L41	12361	SEA ABB=ON	PLU=ON	CHEMICALS/CT
L42	76849	SEA ABB=ON	PLU=ON	L40 OR L41
L43 ·	1869	SEA ABB=ON	PLU=ON	L42 (L) L32
L44	2	SEA ABB=ON	PLU=ON	L43 AND L18
L45	17	SEA ABB=ON	PLU=ON	L39 OR L44
		E SINGH A/A	U	
L46	6030	SEA ABB=ON	PLU=ON	SINGH A?/AU
L47	20767	SEA ABB=ON	PLU=ON	LEE Y?/AU
		E STANISH I	/AU	
L48	22	SEA ABB=ON	PLU=ON	("STANISH I"/AU OR "STANISH IVAN"/AU)
		E DRESSICK	W?/AU	
L49	99	SEA ABB=ON	PLU=ON	DRESSICK W?/AU
		003 300 OM	DT.II-ON	(L46 OR L47 OR L48 OR L49)
L50	26879	SEA ABB=ON	I HOZOM	(210 on 21) on 210 on 21)
L50 L51				L50 AND L18
				· · · · · · · · · · · · · · · · · · ·
	4	SEA ABB=ON D SCAN TI SEA ABB=ON	PLU=ON PLU=ON	L50 AND L18 L50 AND L29
L51	4	SEA ABB=ON D SCAN TI SEA ABB=ON	PLU=ON PLU=ON	L50 AND L18
L51 L52	4 8 95	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON	L50 AND L18 L50 AND L29
L51 L52 L53	4 8 95 2	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16
L51 L52 L53 L54	4 8 95 2 11	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50
L51 L52 L53 L54 L55	4 8 95 2 11 6	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54
L51 L52 L53 L54 L55 L56	4 8 95 2 11 6 7	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54 L53 AND (L19) L53 AND L24 L56 OR L57
L51 L52 L53 L54 L55 L56 L57	4 8 95 2 11 6 7 12	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54 L53 AND (L19) L53 AND L24
L51 L52 L53 L54 L55 L56 L57 L58	4 8 95 2 11 6 7 12 27	SEA ABB=ON D SCAN TI SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON SEA ABB=ON	PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54 L53 AND (L19) L53 AND L24 L56 OR L57 L58 OR L45
L51 L52 L53 L54 L55 L56 L57 L58 L59	4 8 95 2 11 6 7 12 27 6	SEA ABB=ON D SCAN TI SEA ABB=ON	PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54 L53 AND (L19) L53 AND L24 L56 OR L57 L58 OR L45 L55 NOT L59 L53 AND L34
L51 L52 L53 L54 L55 L56 L57 L58 L59 L60	4 8 95 2 11 6 7 12 27 6	SEA ABB=ON D SCAN TI SEA ABB=ON	PLU=ON	L50 AND L18 L50 AND L29 L10 AND L16 L53 AND L50 L51 OR L52 OR L54 L53 AND (L19) L53 AND L24 L56 OR L57 L58 OR L45 L55 NOT L59

=> fil reg
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STRUCTURE FILE UPDATES: 1 NOV 2005 HIGHEST RN 866526-24-1 DICTIONARY FILE UPDATES: 1 NOV 2005 HIGHEST RN 866526-24-1

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REGISTRY includes numerically searchable data for experimental and predicted properties as well as tags indicating availability of experimental property data in the original document. For information on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

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=> d que 15;d 15 rn cn 1-3
L1 ( 1)SEA FILE=REGISTRY ABB=ON PLU=ON "ORGANOPHOSPHORUS HYDROLASE"/
CN
L2 ( 1)SEA FILE=REGISTRY ABB=ON PLU=ON "ORGANOPHOSPHORUS ACID
ANHYDROLASE"/CN
L3 ( 1)SEA FILE=REGISTRY ABB=ON PLU=ON DFPASE/CN
L4 ( 1)SEA FILE=REGISTRY ABB=ON PLU=ON PHOSPHOTRIESTERASE/CN
L5 3 SEA FILE=REGISTRY ABB=ON PLU=ON (L1 OR L2 OR L3 OR L4)
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ANSWER 1 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L5
    117698-12-1 REGISTRY
RN
    Esterase, organophosphate (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
    Aryldialkylphosphatase
CN
    Chlorpyrifos oxonase
CN
CN
    Diazoxonase
    E.C. 3.1.8.1
CN
    Esterase B1
CN
```

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Esterase E3
CN
CN
     Esterase E4
CN
     Esterase, paraoxon
     Esterase, pirimiphos-methyloxon
CN
CN
     Homocysteine thiolactonase
     Homocysteine thiolactone hydrolase
CN
     Isofenphos hydrolase
CN
CN
     Methamidophos-degrading enzyme
     Organophosphate esterase
CN
     Organophosphate hydrolase
CN
     Organophosphate-degrading enzyme
CN
     Organophosphorous hydrolase
CN
     Organophosphorus esterase
CN
     Organophosphorus hydrolase.
CN
     Paraoxon hydrolase
CN
     Paraoxonase
CN
     Paraoxonase 1
CN
     ANSWER 2 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L5
RN
     9047-01-2 REGISTRY
     Esterase, phosphotri- (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
     Phosphorotriester hydrolase
CN
     Phosphotriester hydrolase
CN
     Phosphotriesterase
CN
     ANSWER 3 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L5
     9032-18-2 REGISTRY
RN
     Diisopropylphosphorofluoridase (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
CN
     DFPase
     Dialkylfluorophosphatase
CN
     Diisopropyl phosphorofluoridate hydrolase
CN
     Diisopropylfluorophosphatase
CN
     Diisopropylfluorophosphonate dehalogenase
CN
     E.C. 3.1.8.2
CN
     E.C. 3.8.2.1
CN
     Isopropylphosphorofluoridase
CN
     Organophosphorous acid anhydrolase
CN
     Organophosphorus acid anhydride hydrolase
CN
     Organophosphorus acid anhydrolase
CN
CN
     Sarinase
CN
     Tabunase
=> d que 16 ; d 16 rn cn 1-4
              4 SEA FILE=REGISTRY ABB=ON PLU=ON 9002-98-6 OR 9003-01-4 OR
                49718-51-6 OR 26062-79-3 OR 26062-79-3
     ANSWER 1 OF 4 REGISTRY COPYRIGHT 2005 ACS on STN
L6
     49718-51-6 REGISTRY
RN
     Benzenesulfonic acid, 4-ethenyl-, ion(1-), homopolymer (9CI) (CA INDEX
CN
     NAME)
OTHER NAMES:
CN
     Poly(4-styrenesulfonate)
     ANSWER 2 OF 4 REGISTRY COPYRIGHT 2005 ACS on STN
L6
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26062-79-3 REGISTRY
RN
     2-Propen-1-aminium, N,N-dimethyl-N-2-propenyl-, chloride, homopolymer
CN
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
     Ammonium, diallyldimethyl-, chloride, polymers (8CI)
OTHER NAMES:
CN
     261LV
CN
     Additol VXT 3529
     Agefloc WT 20
CN
     Agefloc WT 20VHV
CN
     Agefloc WT 2206
CN
     Agefloc WT 2206-1800
CN
     Agefloc WT 35VHV
CN
     Agefloc WT 35VLV
CN
     Agefloc WT 40
CN
     Agefloc WT 40HV
CN
     Agefloc WT 40SLV
CN
     Agefloc WT 50SLV
CN
     Agequat 400
CN
     Alcofix 109
CN
     Alcofix 159
CN
     Alcofix 169
CN
     Alcofix 182
CN
     Aronfloc C 70
CN
CN
     AX 04
     AX 05
CN
     AX 05 (polymer)
CN
     Bufloc 536
CN
     Calgon 261
CN
     Calgon 261LV
CN
CN
     Calgon 261RV
     Calgon CP 1030
CN
CN
     Calgon CP 2253
     Calgon CP 261XLV
CN
     Calgon CP 280
CN
     Calgon DMDACC
CN
CN
     Calgon E 904
     Calgon E 905
CN
     Calgon E 921
CN
     Calgon Polymer 261
CN
     Cartafix VXT
CN
CN
     Cat-Floc
     Cat-Floc 71259
CN
CN
    .Cat-Floc L
     Cat-Floc P 112-115
CN
     Cat-Floc T 2
CN
     Cat-Floc TL
CN
CN
     Catiofast CS
CN
     Certrex 340
     CinFix RDF
CN
CN
     CM 100
     CM 100 (onium compound)
CN
CN
     Conductive Polymer 261
CN
     CP 103
     CP 1030
CN
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
     ANSWER 3 OF 4 REGISTRY COPYRIGHT 2005 ACS on STN
Lб
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9003-01-4 REGISTRY

RN

```
2-Propenoic acid, homopolymer (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
    Acrylic acid, polymers (8CI)
OTHER NAMES:
    A 10H
CN
     A 10LL
CN
CN
     A 10SL
     AC 10H
CN
    AC 10LP
CN
     Accusol 440
CN
     Acryl AG 1000
CN
     Acryl AG 1100
CN
     Acryl AG 1200
CN
     Acrylic acid homopolymer
CN
     Acrylic acid polymer
CN
     Acrylic acid resin
CN
     Acrysol A 1
CN
     Acrysol A 3
CN
     Acrysol A 5
CN
CN
     Acrysol AC 5
     Acrysol LMW 20X
CN
     Acrytex W 240
CN
     Acumer 1530
CN
     Acumer 1540
CN
     Acumer 9932
CN
CN
     Acusol 445
     Alcoprint PTF-Z 328
CN
     Alcosperse 409
CN
     Alcosperse 465
CN
     Alcosperse 602A
CN
CN
     AQ 3930
     Aquafeed 600
CN
CN
     Aqualic AS 58
CN
     Aqualic DL 400
     Aqualic HL 321
CN
     Aqualic HL 415
CN
CN
     Aqualic HL 580
CN
     Aquatreat AR 6
CN
     Aquatreat AR 7H
CN
     Arasorb 750
     Arasorb S 100F
CN
CN
     Arespol
CN
     Aron
     Aron 104
CN
CN
     Aron 10H
CN
     Aron A 10H
CN
     Aron A 10LL
     Aron A 10U
CN
CN
     Aron A 30LL
CN
     AS 58
     AS 7503
CN
CN
     ASS 2
CN
     AW 36
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
L6
     ANSWER 4 OF 4 REGISTRY COPYRIGHT 2005 ACS on STN
RN
     9002-98-6 REGISTRY
     Aziridine, homopolymer (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
```

Ethylenimine, polymers (8CI)

```
OTHER NAMES:
CN
     15T
     210T
CN
     2MB
CN
     A 131X
CN
     AC 871
CN
     Adcote 372
CN
CN
     Aziridine polymer
     Basocoll PR 8086
CN
     Basonal White FO 1
CN
     Bufloc 595
CN
     CF 218
CN
     CF 218 (polymer)
CN
     Corcat P 100
CN
     Corcat P 12
CN
     Corcat P 145
CN
     Corcat P 150
CN
     Corcat P 18
CN
CN
     Corcat P 200
     Corcat P 600
CN
     Duramax 1007
CN
     EA 275
CN
     EL 402
CN
     EL 420
CN
CN
     Emerlube 6717
CN
     Epomin
     Epomin 1000
CN
     Epomin 1050
CN
     Epomin 150T .
CN
CN
     Epomin D 3000
     Epomin P 003
CN
CN
     Epomin P 012
     Epomin P 1000
CN
     Epomin P 1030
CN
     Epomin P 1050
CN
     Epomin P 1500
CN
     Epomin P 200
CN
     Epomin P 500
CN
     Epomin SP 003
CN
     Epomin SP 006
CN
     Epomin SP 012
CN
     Epomin SP 018
CN
     Epomin SP 1000
CN
     Epomin SP 103
CN
     Epomin SP 110
CN
     Epomin SP 200
CN
     Epomin SP 300
CN
     Ethyleneimine homopolymer
CN
     Ethylenimine homopolymer
CN
CN
     Everamine
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
  · DISPLAY
=> d que 17;d 17 rn cn 1-3
              3 SEA FILE=REGISTRY ABB=ON PLU=ON 5919-74-4 OR 149305-62-4 OR
                1760-24-3
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ANSWER 1 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L7
     149305-62-4 REGISTRY
RN
     1,2-Propanediol, 3-[(4-ethenylphenyl)methoxy]- (9CI) (CA INDEX NAME)
CN
OTHER NAMES:
     1,2-Dihydroxypropyl 4-vinylbenzyl ether
CN
     ANSWER 2 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L7
     5919-74-4 REGISTRY
RN
     2-Propenoic acid, 2-methyl-, 2,3-dihydroxypropyl ester (9CI) (CA INDEX
CN
     NAME)
OTHER CA INDEX NAMES:
     Methacrylic acid, 2,3-dihydroxypropyl ester (6CI)
     Methacrylin, 1-mono- (8CI)
OTHER NAMES:
     2,3-Dihydroxypropyl methacrylate
CN
     Glyceryl methacrylate
     ANSWER 3 OF 3 REGISTRY COPYRIGHT 2005 ACS on STN
L7
RN
     1760-24-3 REGISTRY
     1,2-Ethanediamine, N-[3-(trimethoxysilyl)propyl]- (9CI) (CA INDEX NAME)
CN
OTHER CA INDEX NAMES:
     Ethylenediamine, N-[3-(trimethoxysilyl)propyl]- (6CI, 7CI, 8CI)
OTHER NAMES:
CN
     (2-Aminoethyl) (3-(trimethoxysilyl)propyl)amine
     (Trimethoxysilylpropyl) ethylenediamine
CN
     \gamma-(2-Aminoethyl)aminopropyltrimethoxysilane
CN
     γ-(Ethylenediamino)propyltrimethoxysilane
CN
     3-(Trimethoxysilyl)propylethylenediamine
CN
CN
     3-[N-(2-Aminoethyl)amino]propyltrimethoxysilane
     6-Amino-4-azahexyltrimethoxysilane
CN
CN
     A 0700
CN
     A 1120
CN
     A 1122
     A 1200
CN
     A 1200 (amine)
CN
CN
     AAS-M
     AO 700
CN
     AP 132
CN
     Dow Corning Z 6020
CN
     Dynasylan DAMO
CN
CN
     Dynasylan DAMO-T
CN
     en-APTAS
CN
     G 91
CN
     G 91 (coupling agent)
CN
     GF 91
CN
     Hydrosil 2776
CN
     K 1600
CN
     KBM 603
CN
     KH 792
CN
     LS 2480
CN
     LS 3750
CN
     N-(\beta-Aminoethyl)-\gamma-aminopropyltrimethoxysilane
     N-(\beta-Aminoethyl)-3-aminopropyltrimethoxysilane
CN
     N-(2-Aminoethyl)-3-aminopropyltrimethoxysilane
CN
CN
     N-(Aminoethyl)aminopropyltrimethoxysilane
CN
     N-(Trimethoxysilylpropyl)ethylenediamine
CN
     N-Aminoethyl-3-aminopropyltrimethoxysilane
CN
     N-[3-(Trimethoxysilyl)propyl]-1,2-ethanediamine
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N-[3-(Trimethoxysilyl)propyl]-1,2-ethylenediamine
CN
    N-[3-(Trimethoxysilyl)propyl]ethylenediamine
CN
    NUCA 1120
CN
     Prosil 3128
CN
     S 320
CN
     SG-SI 900
CN
     SH 6020
CN
     SIA 0591.0
CN
     Sila-Ace S 320
CN
     Silane A 1120
CN
     Silquest A 1120
CN
CN
     SZ 6020
     Trimethoxy[3-[(2-aminoethyl)amino]propyl]silane
CN
CN
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
=> d que 135;d rn cn 135
              1 SEA FILE=REGISTRY ABB=ON PLU=ON COPPER/CN
L35 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2005 ACS on STN
RN
     7440-50-8 REGISTRY
                              (CA INDEX NAME)
     Copper (7CI, 8CI, 9CI)
OTHER NAMES:
CN
     1000Y
CN
     100RXH
CN
     1050Y
CN
     1100T
CN
     1100Y
CN
     115A
CN
     1300YM
CN
     1300YP
CN
     1721 Gold
CN
     200RL
CN
     22BB400
CN
     2L3GT
CN
     3EC
CN
     3EC-HTE
CN
     3EC-III
CN
     3EC-M3S-HTE
CN
     3EC-M3VLP18
CN
     3EC-VEP
CN
     3EC-VLP
CN
     3EC-VLP18
CN
     3EC-VLP35
CN
     3EC3
CN
     3L Fire
CN
     Allbri Natural Copper
CN
     AM-FN
CN
     Arwood copper
CN
     B-WS
CN
     B152-ETP
CN
     BAC 13B-NK120
CN
     BAC 13T
CN
     BHN
CN
     BHN 02T
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BHY 02B-T
CN
     BHY 13B-T
CN
     BHY 13HT
CN
     BHY 13T
CN
     BHY 22B-T
CN
     BHY 22T
CN
     BPF 18
CN
CN
     BSH
     BSH (metal)
CN
     BYH 22B-T
CN
     C 100
CN
     C 100 (metal)
CN
     C.I. 77400
CN
     C.I. Pigment Metal 2
CN
CN
     CCL-HL 830
CN
     CDX
     CDX (metal)
CN
CN
     CE 1100
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
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=> fil_caplus

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FILE COVERS 1907 - 3 Nov 2005 VOL 143 ISS 19
FILE LAST UPDATED: 2 Nov 2005 (20051102/ED)
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Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/infopolicy.html
'OBI' IS DEFAULT SEARCH FIELD FOR 'CAPLUS' FILE

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=> d que 162
               1) SEA FILE=REGISTRY ABB=ON PLU=ON "ORGANOPHOSPHORUS HYDROLASE"/
L1
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                 CN
               1) SEA FILE=REGISTRY ABB=ON PLU=ON "ORGANOPHOSPHORUS ACID
L2
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               1) SEA FILE=REGISTRY ABB=ON PLU=ON PHOSPHOTRIESTERASE/CN 3 SEA FILE=REGISTRY ABB=ON PLU=ON (L1 OR L2 OR L3 OR L4)
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L5
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                 49718-51-6 OR 26062-79-3 OR 26062-79-3
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                 1760-24-3
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L10
         18108 SEA FILE=CAPLUS ABB=ON
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L11
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L12
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L13
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L15
          6408 SEA FILE=CAPLUS ABB=ON PLU=ON L12 OR L13
L16
             2 SEA FILE=CAPLUS ABB=ON PLU=ON L14 AND L15 AND L16
L17
           146 SEA FILE=CAPLUS ABB=ON PLU=ON L14 AND L15
L18
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L19
       2314372 SEA FILE=CAPLUS ABB=ON PLU=ON MATERIAL/OBI OR BIOACTIV?/OBI
L21
               OR GLASS/OBI OR SILICA/OBI OR TEXTILE#/OBI OR CLOTH?/OBI OR
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L25
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L27
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L29
           160 SEA FILE=CAPLUS ABB=ON PLU=ON L21 (L) L29
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            17 SEA FILE=CAPLUS ABB=ON PLU=ON L30 AND L19
L31
        950582 SEA FILE=CAPLUS ABB=ON PLU=ON PROTECT?/OBI OR DECONTAMIN?/OBI
L32
                OR SAFETY/OBI OR COAT?/OBI
             4 SEA FILE=CAPLUS ABB=ON PLU=ON L31 AND L32
L33
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L39
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                                      PLU=ON CHEMICALS/CT
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         76849 SEA FILE=CAPLUS ABB=ON
L42
                                              L42 (L) L32
          1869 SEA FILE=CAPLUS ABB=ON
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L43
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                                              L10 AND L16
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L57
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L58
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                                       PLU=ON
                                              L53 AND L34
L61
                                       PLU=ON L61 OR L59
             28_SEA_FILE=CAPLUS_ABB=ON
L62
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1760-24-3

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L8
         6210 SEA FILE=CAPLUS ABB=ON PLU=ON ENZYMES, USES/CT
L9
        31031 SEA FILE=CAPLUS ABB=ON PLU=ON L6
L10
        18108 SEA FILE=CAPLUS ABB=ON PLU=ON POLYELECTROLYTES/CT
L11
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         46227 SEA FILE=CAPLUS ABB=ON PLU=ON L10 OR L11
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T<sub>2</sub>1
               OR GLASS/OBI OR SILICA/OBI OR TEXTILE#/OBI OR CLOTH?/OBI OR
               THREAD?/OBI
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L22
.L23
        10530 SEA FILE=CAPLUS ABB=ON PLU=ON DECONTAMI?/OBI
        737361 SEA FILE=CAPLUS ABB=ON PLU=ON CATALYS?/OBI
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L25
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L27
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L29
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L33
L34
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L35
L36
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L37
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L39
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L41
          76849 SEA FILE=CAPLUS ABB=ON PLU=ON L40 OR L41
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          6030 SEA FILE=CAPLUS ABB=ON PLU=ON SINGH A?/AU
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          20767 SEA FILE=CAPLUS ABB=ON PLU=ON LEE Y?/AU
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             22 SEA FILE=CAPLUS ABB=ON PLU=ON ("STANISH I"/AU OR "STANISH
L48
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             99 SEA FILE=CAPLUS ABB=ON PLU=ON DRESSICK W?/AU
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L50
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                                      PLU=ON L58 OR L45
           27 SEA FILE=CAPLUS ABB=ON
L59
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1 SEA FILE=CAPLUS ABB=ON PLU=ON L53 AND L34
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L61
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                                              L61 OR L59
L62
        6_SEA_FILE=CAPLUS ABB=ON
                                       PLU=ON
                                               L60 NOT L62
L63
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=> d .ca 162 1-28;d .ca 163 1-6 THE ESTIMATED COST FOR THIS REQUEST IS 83.16 U.S. DOLLARS DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N: γ

L62 ANSWER 1 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2005:641826 CAPLUS

DOCUMENT NUMBER: 143:156038

TITLE: Methods of using sealants in multilateral junctions

INVENTOR(S): Eoff, Larry S.; Everett, Don M.

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 14 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005159319	A1	20050721	US 2004-759676	20040116
PRIORITY APPLN. INFO.:			US 2004-759676	20040116

ED Entered STN: 22 Jul 2005

- The present invention relates to the completion of subterranean well bores in a multilateral well system. More particularly, this invention relates to the sealing of junctions between lateral well bores and a parent well bore. The sealants used in accordance with the methods of the present invention generally comprise any sealing composition which can be placed within a reservoir, and injected a sufficient distance into a region of the formation surrounding a junction between a 1st well bore and a 2nd well bore in fluid communication with the 1st well bore, so as to prevent the undesired entry of formation fluids into either well bore in the region surrounding the junction.
- IC ICM E21B033-00
- INCL 507225000; 507219000; 507224000
- CC 51-2 (Fossil Fuels, Derivatives, and Related Products) Section cross-reference(s): 38, 39
- IT Polymerization catalysts
 - (delayed activator for silicates; methods of using polymer and copolymer sealants in multilateral junctions)
- IT Polymerization catalysts
 - (radical; methods of using polymer and copolymer sealants in multilateral junctions)
- IT Polymerization catalysts
 - (redox; methods of using polymer and copolymer sealants in multilateral junctions)
- IT 919-30-2, γ -Aminopropyltriethoxysilane 1760-24-3
 - 3069-24-7 35141-30-1 103526-27-8 860028-10-0
 - RL: MOA (Modifier or additive use); USES (Uses)
 - (coupling agent; methods of using polymer and copolymer sealants in multilateral junctions)
- IT 9002-98-6
 - RL: RCT (Reactant); TEM (Technical or engineered material use); RACT (Reactant or reagent); USES (Uses)
 - (crosslinking agent; methods of using polymer and copolymer sealants in multilateral junctions)
- IT 64-18-6, Formic acid, uses 64-19-7, Acetic acid, uses 76-03-9, Trichloroacetic acid, uses 79-52-7, 1,1,3-Trichlorotrifluoroacetone 85-42-7, Hexahydrophthalic anhydride 85-44-9, Phthalic anhydride 88-95-9, 1,2-Benzenedicarbonyl dichloride 89-32-7, Pyromellitic dianhydride 98-07-7, Benzotrichloride 98-87-3, Benzal chloride

98-88-4, Benzoyl chloride 100-44-7, Benzyl chloride, uses Diaminodiphenylmethane 103-83-3, Benzyldimethylamine 104-78-9 107-15-3, Ethylenediamine, uses 108-31-6, Maleic anhydride, uses 108-45-2, m-Phenylenediamine, uses 109-55-7 110-89-4, Piperidine, uses 111-40-0, Diethylenetriamine 112-24-3, Triethylenetetramine Hexachloroacetone 140-31-8, 1-Piperazineethanamine 627-63-4, Fumaryl 694-83-7, 1,2-Diaminocyclohexane 1477-55-0. chloride 1,3-Benzenedimethanamine 2855-13-2, Isophorone diamine 7647-01-0. Hydrochloric acid, uses 9002-98-6D, derivs. 26444-72-4, (Tris(dimethylamino)methyl)phenol 26590-20-5, Methyltetrahydrophthalic anhydride 28299-33-4, Imidazoline 31307-24-1, Methylbicyclo-[2,2,1]-5heptene-2,3-dicarboxylic anhydride 59516-66-4, Oxalic anhydride 860309-87-1 RL: CAT (Catalyst use); RCT (Reactant); RACT (Reactant or reagent); USES (Uses)

(hardening agent; methods of using polymer and copolymer sealants in multilateral junctions)

L62 ANSWER 2 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

2005:547065 CAPLUS

DOCUMENT NUMBER:

143:73874

TITLE:

Catalytic enzyme-modified

textiles for active protection from

toxins

INVENTOR(S):

Singh, Alok; Dressick, Walter J.; Lee, Yongwoo

USA

SOURCE: -

U.S. Pat. Appl. Publ., 18 pp., Sont.-in-part of U.S.

Ser. No. 750,637.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

ED

AB

English

FAMILY ACC. NUM. COUNT: 2

Entered STN: 24 Jun 2005

PATENT INFORMATION:

PATENT ASSIGNEE(S):

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005136523	A1	20050623	US 2004-849621	20040520
US 2005136522	A1	20050623	US 2003-750637	20031223
PRIORITY APPLN. INFO.:			US 2003-750637	A2 20031223

The present invention provides bioactive catalytic enzyme-modified textiles for providing protection from chemical exposure that are stable and retain their catalytic activity comprise a cloth substrate, at least one polyelectrolyte layer, at least one enzyme layer to degrade the chemical agent, and an end-capping layer. The textiles of the present invention can be used for reusable clothing that decontaminates itself after exposure to toxins and can be worn multiple times or for disposable clothing and wipes intended for a single use that decontaminate themselves without harming the environment. In a preferred embodiment, the present invention takes advantage of superior catalytic activity of enzymes by immobilizing them within polyelectrolyte multilayers (PEMs). technique for forming multilayers is simple and effective as polyelectrolytes of opposing polarity are alternatively deposited through neutralization and overcompensation of their charges. A capping agent provides stability to the multilayers, keeps enzymes protected in adverse working environments, and attracts the toxic agents to facilitate contact with the catalytic sites. The present invention provides several advantages over the prior art. It leads to enhanced enzyme shelf life under normal storage conditions. It allows incorporation of multiple

components into multilayers to provide add- on capabilities to the

```
packaged system. It is lightwt., robust, sturdy, disposable,
     self-decontaminating, and cost-effective. It offers versatility as it can
     be designed for use on various materials. One example demonstrates the
     preparation of a polyelectrolyte-enzyme multilayer containing a single layer
of the
     OPH enzyme on a woven fiberglass cloth sample by the dip coating method.
     ICM C12N009-16
TC
INCL 435196000
     7-7 (Enzymes)
CC
     Section cross-reference(s): 40
     catalysis enzyme modified textile
ST
     toxin decontamination protective
     clothing
IT
     Chemicals
       Decontamination
       Textiles
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Enzymes, uses
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Fluoropolymers, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Toxins
     RL: REM (Removal or disposal); PROC (Process)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Glass fiber fabrics
     RL: TEM (Technical or engineered material use); USES (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
     Glass fibers, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
     Polyamide fibers, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Rayon, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (catalytic enzyme-modified textiles for
        active protection from toxins)
IT
     Polymers, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (charged, substrate capable of adsorbing; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     Safety devices
IT
        (chemical protective clothing; catalytic
        enzyme-modified textiles for active
        protection from toxins)
IT
     Clothing
        (chemical protective; catalytic enzyme-
        modified textiles for active protection
        from toxins)
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Amines, uses
TT
    RL: NUU (Other use, unclassified); USES (Uses)
        (cotton modified with; catalytic enzyme-
        modified textiles for active protection
        from toxins)
    Textiles
TΤ
      Textiles
        (cotton; catalytic enzyme-modified textiles
        for active protection from toxins)
     Coating process
IT
        (dip, layers are deposited using; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     Functional groups
        (hexyl, outmost capping layer comprises branched polyethylenimine
        (BPEI) modified with; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     Polyelectrolytes
IT
        (layer; catalytic enzyme-modified textiles
        for active protection from toxins)
     Antibacterial agents
TΤ
        (outermost capping layer contains; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     Acrylic polymers, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises; catalytic enzyme-
        modified textiles for active protection
        from toxins)
ΙT
     Coating process
        (spin, layers are deposited using; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     Coating process
        (spray, layers are deposited using; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     Functional groups
        (substrate comprises a material whose surface chemical
        modified to generate; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     Threads
        (woven into a fabric, substrate is; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     923-26-2, 1,2-Dihydroxypropyl methacrylate 1760-24-3,
IT
     N-[(3-Trimethoxysilyl)propyl]ethylenediamine 149305-62-4,
     1,2-Dihydroxypropyl 4-vinylbenzyl ether
     RL: NUU (Other use, unclassified); USES (Uses)
        (branched, capping layer comprises; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     12619-70-4, Cyclodextrin
     RL: NUU (Other use, unclassified); USES (Uses)
        (cotton modified with; catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     7585-39-9, β-Cyclodextrin
```

```
RL: NUU (Other use, unclassified); USES (Uses)
        (functionalized polyelectrolyte, polyelectrolyte layer comprises;
        catalytic enzyme-modified textiles for
        active protection from toxins)
    9032-18-2, Organophosphorous acid anhydrolase
TT
    RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
        (organophosphorous acid anhydrolase; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     117698-12-1, EC 3.1.8.1
IT
    RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (organophosphorous hydrolase; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     74-83-9, Methyl bromide, uses
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (outmost capping layer comprises branched polyethylenimine quaternized
        with; catalytic enzyme-modified textiles
        for active protection from toxins)
     9002-84-0, Polytetrafluoroethylene
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (oxidized, substrate comprises; catalytic enzyme-
        modified textiles for active protection
        from toxins)
     9047-01-2, Phosphotriesterase
IT
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (phosphotriesterases (PTE) (EC 3.1.8); catalytic enzyme-
        modified textiles for active protection
        from toxins)
IT
     50851-57-5
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises, capping layer comprises; catalytic
        enzyme-modified textiles for active
        protection from toxins)
                9003-47-8, Polyvinylpyridine
IT
     9002-98-6
                                                25087-26-7,
                            25191-25-7, Polyvinyl sulfate 26062-79-3,
     Polymethacrylic acid
                                            26913-06-4, Poly[imino(1,2-
     Polydiallyl dimethylammonium chloride
                   71550-12-4, Polyallylamine hydrochloride
     ethanediyl)]
     RL: NUU (Other use, unclassified); USES (Uses)
        (polyelectrolyte layer comprises; catalytic enzyme-
        modified textiles for active protection
        from toxins)
L62 ANSWER 3 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                         2005:547064 CAPLUS
DOCUMENT NUMBER:
                         143:82626
                         Catalytic surfaces for active protection from
TITLE:
                         toxins
                         Singh, Alok; Lee, Yongwoo; Stanish, Ivan; Chang,
INVENTOR(S):
                         Eddie; Dressick, Walter J.
PATENT ASSIGNEE(S):
                         U.S. Pat. Appl. Publ., 15 pp.
SOURCE:
                         CODEN: USXXCO
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
```

	PATENT NO.	KIND	DATE	APPLICATION NO.	
	US 2005136522		20050623		
	US 2005136523		20050623	US 2004-849621	20040520
PRTO	RITY APPLN. INFO.:	***	20030023	US 2004-849621 US 2003-750637	A2 20031223
ED	Entered STN: 24 Ju	n 2005			
AB	The present inventi		tes to cata	lytic surfaces for	active protection
	from air or water b				
	materials. A bioac	tive ca	talytic mat	erial is disclosed	for providing this
	protection from che	mical e	xposure. T	he material is com	posed of enzymes
	immobilized within	polyele	ctrolyte mu	ltilayers and a po	lymerizable
	end-capping layer t	o rende	er stability	to enzymes. Also	disclosed is the
	related method for				
	deposition on subst				xibility for
	providing active pr	cotection	on from chem	nical exposure.	
IC	ICM A61K038-46				
	ICS C12N009-16				
	435196000; 42409460		. J	:)	
CC	59-5 (Air Pollution Section cross-refer			giene)	
ST	multilayer bioactiv	ence(s)	: 4 Vtic materi	al towin deardn ad	sorption:
51	chem warfare agent	toxin d	lycic maceri Begrdn adsor	ntion bioactive	sorption,
	catalytic material;	protec	tive clothi	ng chem warfare ag	ent
	bioactive catalytic			3	
IT	Toxins				
	RL: ADV (Adverse ef	fect, i	ncluding to	oxicity); PEP (Phys	ical, engineering
	or chemical process				val or disposal);
	BIOL (Biological st				
		orne; ca	talytic sur	faces for active p	rotection
	from toxins)				
IT	Glass beads Glass fiber fabrics				
	RL: TEM (Technical		neered mate	rial use) · HSES (I	Ises)
	(as substrate: (ratalvti	c surfaces	for active protect	ion from
	toxins)	, u o u = 1 - 1		Parties	
IT	Adsorption				
	Chemical warfare	agents			
	Hydrolysis kinetics				
	Passivation				
	-	aces for	c active pro	otection from toxi n	s)
IT	Enzymes, uses			tal and more than	· \
	RL: TEM (Technical				
~~	(catalytic surfa	ices Ioi	active pro	tection from toxin	is)
IT		trate.	catalytic s	surfaces for active	protection from
	toxins)	scrace,	catalytic	dilaces for accive	proceeding from
IT	Hydrolysis			•	
		alytic s	surfaces for	active protection	from
	toxins)	•		•	
IT	Safety devices				
	(protective clot	hing; o	catalytic su	rfaces for active	protection from
	toxins)				
IT	Clothing				
		calytic	surfaces fo	or active protection	on from
	toxins)				
ΙT	Humidity			المال المستعدين المراوع والمراوع والمرا	auxfacca for acti
				ctivity; catalytic	surfaces for active
IT	protection from 298-00-0, Methylpan			•	
11	296-00-0, Methylpa	Lacinton		•	

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RL: ADV (Adverse effect, including toxicity); CPS (Chemical process); PEP
     (Physical, engineering or chemical process); REM (Removal or disposal);
     BIOL (Biological study); PROC (Process)
        (catalytic surfaces for active protection from toxins)
     74173-48-1D, 4-Vinyl-4'-methyl-2,2'-bipyridine, copper complexes, polymers
TΤ
    with trimethylolpropane trimethacrylate
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (catalytic surfaces for active protection from toxins)
     3290-92-4D, Trimethylolpropane trimethacrylate, polymer with Cu(II)-containing
IT
     vinyl compds. 9032-18-2, Organophosphorus acid anhydrolase
     9047-01-2, Phosphotriesterase 117698-12-1,
     Organophosphorus hydrolase
     RL: TEM (Technical or engineered material use); USES (Uses)
        (catalytic surfaces for active protection from toxins)
     15158-11-9D, complexes, contained in vinyl compds., polymer with
IT
     trimethylolpropane trimethacrylate
     RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES
     (Uses)
        (complex with ethanediamidoxime; catalytic surfaces for active
        protection from toxins)
     79647-56-6, Poly-β-cyclodextrin
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (crosslinked, for adsorption of paranitrophenol; catalytic surfaces for
        active protection from toxins)
     7647-14-5, Sodium chloride, miscellaneous
IT
     RL: MSC (Miscellaneous)
        (effect of, on enzyme activity; catalytic surfaces for active
        protection from toxins)
     1760-24-3, N-[3-Trimethoxysilyl)propyl]ethylenediamine
IT
     5919-74-4 149305-62-4
     RL: TEM (Technical or engineered material use); USES (Uses)
        (end-capping agent; catalytic surfaces for active
        protection from toxins)
     55-91-4, Diisopropyl fluorophosphate
IT
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); REM (Removal or disposal); PROC (Process)
        (nerve agent simulant; catalytic surfaces for active protection from
        toxins)
IT
     7631-86-9, Silica, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (of glass beads, as substrate; catalytic surfaces for active protection
        from toxins)
     9002-98-6 9003-01-4, Polyacrylic acid 26062-79-3
IT
     , Polydiallyl dimethyl ammonium chloride 50851-57-5
     RL: TEM (Technical or engineered material use); USES (Uses)
        (or its derivs. branched or linear; catalytic surfaces for active
        protection from toxins)
                          1112-38-5
IT
     100-02-7, processes
     RL: CPS (Chemical process); FMU (Formation, unclassified); PEP (Physical,
     engineering or chemical process); FORM (Formation, nonpreparative); PROC
     (Process)
        (product of methylparathion degradation; catalytic surfaces for active
        protection from toxins)
L62 ANSWER 4 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                         2005:239795 CAPLUS
ACCESSION NUMBER:
TITLE:
                         Self-cleaning fabrics for decontamination of
                         organophosphorus pesticides and related chemical
                         agents. [Erratum to document cited in CA142:256107]
```

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Singh, Alok; Lee, Yongwoo; Dressick, Walter J.
AUTHOR (S):
                         Center for Bio/Molecular Science and Engineering,
CORPORATE SOURCE:
                         Naval Research Laboratory, Washington, DC, 20375, USA
SOURCE:
                         Advanced Materials (Weinheim, Germany) (2005), 17(4),
                         CODEN: ADVMEW; ISSN: 0935-9648
PUBLISHER:
                         Wiley-VCH Verlag GmbH & Co. KGaA
                         Journal; Errata
DOCUMENT TYPE:
                         English
LANGUAGE:
     Entered STN: 18 Mar 2005
ED
AB
     An erratum.
     4-4 (Toxicology)
CC
     Section cross-reference(s): 5
     erratum self cleaning fabric decontamination organophosphorus
ST
     pesticide
     INDEXING IN PROGRESS
IT
     Textiles
TT
        (cotton; on self-cleaning fabrics for decontamination of
        organophosphorus pesticides (Erratum))
     Decontamination
IT
        (on self-cleaning fabrics for decontamination of
        organophosphorus pesticides (Erratum))
TΤ
     Pesticides
        (organophosphorus; on self-cleaning fabrics for decontamination
        of organophosphorus pesticides (Erratum))
     Glass fibers
IT
     RL: MSC (Miscellaneous)
        (woven cloth; on self-cleaning fabrics for
        decontamination of organophosphorus pesticides (Erratum))
IT
     117698-12-1, Organophosphorus hydrolase
     RL: BUU (Biological use, unclassified); BIOL (Biological study); USES
     (Uses)
        (on self-cleaning fabrics for decontamination of
        organophosphorus pesticides (Erratum))
     298-00-0, Methyl parathion
IT
     RL: REM (Removal or disposal); PROC (Process)
        (on self-cleaning fabrics for decontamination of
        organophosphorus pesticides (Erratum))
     9002-98-6, Polyethylenimine
IT
     RL: MOA (Modifier or additive use); USES (Uses)
        (stabilizer; on self-cleaning fabrics for decontamination of
        organophosphorus pesticides (Erratum))
L62 ANSWER 5 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                         2005:59598 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         142:256107
                         Self-cleaning fabrics for decontamination of
TITLE:
                         organophosphorus pesticides and related chemical
                         agents
                       · Singh, Alok; Lee, Yongwoo; Dressick, Walter J.
AUTHOR(S):
                         Center for Bio/Molecular Science and Engineering,
CORPORATE SOURCE:
                         Naval Research Laboratory, Washington, DC, 20375, USA
                         Advanced Materials (Weinheim, Germany) (2004),
SOURCE:
                         16(23-24), 2112-2115
                         CODEN: ADVMEW; ISSN: 0935-9648
                         Wiley-VCH Verlag GmbH & Co. KGaA
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Entered STN: 21 Jan 2005
     Fabrics are coated with polyelectrolyte multilayers containing
AB
```

organophosphorus hydrolase. The fabrics are usable for self-decontamination of methyl parathion in the environment. The authors sequentially deposited an aqueous branched polyethylenimine solution, a organophosphorus hydrolase solution in a buffer and a branched polyethylenimine solution in a buffer on a woven fiberglass cloth, to obtain the title fabric. In another experiment, a cotton fabric was used as a substrate. The self-cleaning fabrics showed high enzyme activity and good durability.

CC 4-4 (Toxicology)

Section cross-reference(s): 5

ST self cleaning fabric decontamination organophosphorus pesticide

IT Textiles

(cotton; on self-cleaning fabrics for **decontamination** of organophosphorus pesticides)

IT Decontamination

(on self-cleaning fabrics for **decontamination** of organophosphorus pesticides)

IT Pesticides

(organophosphorus; on self-cleaning fabrics for **decontamination** of organophosphorus pesticides)

IT Glass fibers, miscellaneous

RL: MSC (Miscellaneous)

(woven cloth; on self-cleaning fabrics for decontamination of organophosphorus pesticides)

IT 117698-12-1, Organophosphorus hydrolase

RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)

(on self-cleaning fabrics for **decontamination** of organophosphorus pesticides)

IT 298-00-0, Methyl parathion

RL: REM (Removal or disposal); PROC (Process)
(on self-cleaning fabrics for decontamination of

organophosphorus pesticides)

Organophosphorus peserere

IT 9002-98-6, Polyethylenimine

RL: MOA (Modifier or additive use); USES (Uses)

(stabilizer; on self-cleaning fabrics for **decontamination** of organophosphorus pesticides)

REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L62 ANSWER 6 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

2005:30978 CAPLUS

DOCUMENT NUMBER:

142:74718

TITLE:

Process for preparation of propyltrichlorosilane by hydrosilylation of propene with trichlorosilane in

presence of supported platinum catalyst with

polymeric membrane

INVENTOR(S):

Marciniec, Bogdan; Foltynowicz, Zenon; Checinski,

Dariusz

PATENT ASSIGNEE(S):

Uniwersytet im. Adama Mickiewicza, Pol.

SOURCE:

Pol., 6 pp. CODEN: POXXA7

DOCUMENT TYPE: Patent LANGUAGE: Polish

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PL 185355	В1	20030430	PL 1997-322149	19970915

```
19970915
PRIORITY APPLN. INFO.:
                                             PL 1997-322149
                         CASREACT 142:74718
OTHER SOURCE(S):
     Entered STN: 14 Jan 2005
     Propyltrichlorosilane is prepared by addition reaction of trichlorosilane to
AΒ
     propene in presence of a platinum catalyst such that the platinum catalyst
     used has a polymeric membrane prepared by condensation reaction of
     methacrylic or acrylic acid, either with a polymerization initiator or
directly,
     with amino groups on the surface of silica modified by silane on which
     there is also a platinum complex prepared by immobilization of
     hexachloroplatinic acid; the reaction takes place at normal pressure in
     the gas phase at 357-372° with a reagent feed rate through the
     catalyst of 1.5-4.5 L/h and a volume ratio of propene to trichlorosilane in
     the reaction mixture of 1.5-2.5. In an example, silica gel
     (MN-Kieselgel-60) is modified by sequential treatments with
     (3-chloropropyl)triethoxysilane, hexamethyldisilazane, sodium
     diethyldithiocarbamate (polymerization initiator), hexachloroplatinic acid, and
     then photochem. polym. of methacrylic acid to form the polymeric membrane
     afforded a catalyst which was then used in the gas-phase hydrosilylation
     of propene at 372 K with trichlorosilane with a reagent feed rate of 1.65
     L/h and a reagent volume ratio of 2.0 gave 69-75% conversion of
     trichlorosilane with a selectivity for propyltrichlorosilane of 100%.
IC
     ICM C07F007-14
     ICS B01J031-28
     29-6 (Organometallic and Organometalloidal Compounds)
CC
     Section cross-reference(s): 35, 45
     process propyltrichlorosilane prepn supported catalyst; propene
st
     hydrosilylation silica supported platinum catalyst polymeric
     membrane process
IT
     Hydrosilylation catalysts
        (platinum; preparation of propyltrichlorosilane by hydrosilylation of
        propene with trichlorosilane in presence of modified silica-supported
        platinum catalyst with polymeric membrane)
     Silica gel, preparation
IT
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (reaction products; preparation of propyltrichlorosilane by hydrosilylation
        of propene with trichlorosilane in presence of modified
        silica-supported platinum catalyst with polymeric membrane)
     148-18-5DP, Sodium diethyldithiocarbamate, modified silica-bound
TΤ
     999-97-3DP, Hexamethyldisilazane, modified silica-bound
     1760-24-3DP, modified silica-bound
                                          5089-70-3DP,
     (3-Chloropropyl) triethoxysilane, modified silica-bound 9003-01-4DP
     , Poly(acrylic acid), modified silica-bound 16941-12-1DP,
     Hexachloroplatinic acid, silica gel- and polymeric membrane-bound 25087-26-7DP, Poly(methacrylic acid), modified silica-bound
     RL: CAT (Catalyst use); SPN (Synthetic preparation); PREP (Preparation);
     USES (Uses)
        (preparation of propyltrichlorosilane by hydrosilylation of propene with
        trichlorosilane in presence of modified silica-supported platinum
        catalyst with polymeric membrane)
     141-57-1P, Propyltrichlorosilane
IT
     RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP
     (Preparation)
        (preparation of propyltrichlorosilane by hydrosilylation of propene with
        trichlorosilane in presence of modified silica-supported platinum
        catalyst with polymeric membrane)
IT
     115-07-1, Propene, reactions
                                    10025-78-2, Trichlorosilane
     RL: RCT (Reactant); RACT (Reactant or reagent)
```

(preparation of propyltrichlorosilane by hydrosilylation of propene with

trichlorosilane in presence of modified silica-supported platinum catalyst with polymeric membrane)

148-18-5, Sodium diethyldithiocarbamate 999-97-3, Hexamethyldisilazane 1760-24-3 5089-70-3, (3-Chloropropyl)triethoxysilane 16941-12-1, Hexachloroplatinic acid

RL: RCT (Reactant); RACT (Reactant or reagent)

(reaction with modified silica surface; preparation of propyltrichlorosilane by hydrosilylation of propene with trichlorosilane in presence of modified silica-supported platinum **catalyst** with polymeric membrane)

L62 ANSWER 7 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2004:780930 CAPLUS

DOCUMENT NUMBER: 141:273966

TITLE: The detecting method and device of polypeptide, and

the ligand compound comprising nanoparticles

INVENTOR(S): Zou, Fanglin; Chen, Chunsheng; Chen, Ning; Wang,

Jianxia

PATENT ASSIGNEE(S): Chengdu Kuachang Medical Industrial Limited, Peop.

Rep. China; Chengdu Kuachang Science & Technology Co.,

Ltd

SOURCE: PCT Int. Appl., 61 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent Chinese

LANGUAGE: C FAMILY ACC. NUM. COUNT: 7

PATENT INFORMATION:

PATENT	NO.			KIN	D :	DATE		,	APPL	ICAT	ION 1	NO.		D	ATE		
		,			-									-			
WO 2004	0815	71		A1	;	2004	0923		WO 2	004-	CN77			2	0040	120	
W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	ВA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	
	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,	
	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
											VC,						
RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	
	BY,	KG,	KΖ,	MD,	RU,	TJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	
	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG
CN 1434	295			Α		2003	0806		CN 2	003-	1174	46		2	0030	313	
CN 1514	243			Α	:	2004	0721		CN 2	003-	1177	87		2	00304	430	
WO 2004	0905	48		A1	;	2004	1021		WO 2	004-	CN20	3		2	0040	315	
W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KΡ,	KR,	ΚZ,	LC,	
	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	ΜZ,	NA,	NI,	
	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	
	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
RW:	BW,	GH,	GM,	KE,	LS,	MW,	ΜZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	
	BY,	KG,	ΚZ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	
	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	
	TD,	TG															
WO 2004	1021	96		A1		2004	1125		WO 2	004-	CN43'	7		2	0040	430	
₩:	ΑE,	AG,	ΑL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FΙ,	GB,	GD,	
	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KZ,	LC,	
	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,	

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NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
             TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM,
             AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
             EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE,
             SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
             SN, TD, TG
PRIORITY APPLN. INFO.:
                                            CN 2003-117446
                                                                A 20030313
                                            CN 2003-117787
                                                                A 20030430
                                                                A 20040120
                                            WO 2004-CN77
                                            WO 2004-CN203
                                                                A 20040315
    Entered STN: 24 Sep 2004
ED
    The invention relates to a quant. and/or qual. detecting method of
AB
    polypeptides, where the objects react with affinity nanocarriers and / or
     ligands / nanoparticles marked mol. compds. resp. The invention also
     relates to the detecting device which relates to affinity nanocarriers and
     / or ligands / nanoparticles / marked mol. compds., especially relates to
chips,
     enzyme marked plates, chromatog. bands. Moreover, the invention relates
     to affinity nanocarriers used as detecting and isolating medium, and the
     method of preparing such carriers. Also, the invention relates to ligands /
     nanoparticles / marked mol. compds. and the method of preparing such compds.
     The invention provides chips, kits, detecting method and device which
     comprise magnetic nanoparticles.
IC
     ICM G01N033-543
     ICS C07K002-00; C12Q001-00
     9-1 (Biochemical Methods)
CC
    Section cross-reference(s): 10, 14, 15
    Analytical apparatus
IT
    Antibiotics
     Biochips
     Blood analysis
     Cell
     Ceramics
     Chemiluminescent substances
     Chromatography
    Dyes
     Fluorescent indicators
    Hepatitis B virus
    Hepatitis C virus
    Molecules
    Nanoparticles
     Plates
       Polyelectrolytes
     Reaction
     Stains, coloring materials
     Surfactants
     Test kits
     Thin layer chromatographs
     Virus
        (detecting method and device of polypeptide, and the ligand compound
        comprising nanoparticles)
IT
    Avidins
       Enzymes, uses
     Ligands
     Polysaccharides, uses
     Salts, uses
     Vitamins
     RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
        (detecting method and device of polypeptide, and the ligand compound
```

comprising nanoparticles)

IT 1344-28-1, Aluminum oxide, analysis 7429-90-5, Aluminum, analysis

7440-21-3, Silicon, analysis 7440-22-4, Silver, analysis

7440-50-8, Copper, analysis 7440-57-5, Gold, analysis

7631-86-9, Silica, analysis 13463-67-7, Titanium oxide, analysis

RL: ARU (Analytical role, unclassified); ANST (Analytical study)

(detecting method and device of polypeptide, and the ligand compound

comprising nanoparticles)

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L62 ANSWER 8 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2004:757067 CAPLUS

DOCUMENT NUMBER: 141:246119

TITLE: Biocatalytic electrode with switchable and tunable

power output and fuel cell using such electrode

INVENTOR(S): Katz, Eugenii; Willner, Itamar

PATENT ASSIGNEE(S): Yissum Research Development Company of the Hebrew

University of Jerusalem, Israel

SOURCE: PCT Int. Appl., 44 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

REFERENCE COUNT:

PAT	ENT NO.				KIND DATE			APPLICATION NO.						DATE			
		- -				-											
WO	2004	07984	18		A2		2004	916	1	WO 2	004-3	[L19	9		20040302		
WO	2004	07984	48		A 3		2005	0506									
	W:	ΑE,	ΑE,	AG,	AL,	AL,	AM,	AM,	AM,	AT,	AT,	AU,	ΑZ,	ΑZ,	BA,	BB,	BG,
		BG,	BR,	BR,	BW,	BY,	BY,	ΒZ,	ΒZ,	CA,	CH,	CN,	CN,	CO,	CO,	CR,	CR,
		CU,	CU,	CZ,	CZ,	DE,	DE,	DK,	DK,	DM,	DZ,	EC,	EC,	EE,	EE,	EG,	ES,
		ES,	FI,	FI,	GB,	GD,	GE,	GE,	GH,	GM,	HR,	HR,	HU,	HU,	ID,	IL,	IN,
		IS,	JP,	JP,	KE,	ΚE,	KG,	KG,	ΚP,	KP,	KP,	KR,	KR,	KZ,	ΚZ,	ΚZ,	LC,
		LK,	LR,	LS,	LS,	LT,	LU,	LV,	MA,	MD,	MD,	MG,	MK,	MN,	MW,	MX,	MX,
		MZ,	ΜZ,	NA,	NI												
	RW:	BW,	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	ΑT,	ΒE,
		BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	ΙΤ,	LU,
		MC,	ΝL,	ΡL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,
		GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,
		GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG							
PRIORITY	APPLN. INFO.:			. :				•	, 1	US 2	003-	4507	02P	1	P 20	0030	303

ED Entered STN: 16 Sep 2004

The present invention provides a novel electrode carrying on at least a portion of its support surface a hybrid polymer matrix (HPM), a catalyst that can catalyze a redox reaction and an optional electron mediator group that enhances the elec. contact between the HPM and the catalyst, the HPM being capable to be electrochem. changed from a non-conductive state to a conductive state. The electrode of the invention may be used in elec. devices such as fuel cells, thus imparting them switchable and tunable properties. The fuel cell of the invention may be used as a power source or as a self-powered sensor.

- IC ICM H01M008-16
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 9, 38
- IT Enzymes, uses

RL: CAT (Catalyst use); USES (Uses)

(redox; biocatalytic electrode with switchable and tunable power output and fuel cell using such electrode)

TT 7439-89-6, Iron, uses 7439-97-6, Mercury, uses 7440-02-0, Nickel, uses
7440-22-4, Silver, uses 7440-47-3, Chromium, uses 7440-50-8,
Copper, uses 7440-57-5, Gold, uses 7440-66-6, Zinc, uses
9003-01-4, Polyacrylic acid 25104-18-1, Polylysine 50851-57-5
RL: DEV (Device component use); USES (Uses)
 (biocatalytic electrode with switchable and tunable power output and fuel cell using such electrode)

L62 ANSWER 9 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2004:433703 CAPLUS

DOCUMENT NUMBER: 141:9611

TITLE: Enzyme immobilization for use in biofuel cells and

sensors

INVENTOR(S): Minteer, Shelley D.; Akers, Niki L.; Moore, Christine

Μ.

PATENT ASSIGNEE(S): St. Louis University, USA

SOURCE: U.S. Pat. Appl. Publ., 33 pp., which

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.					KIND DATE		APPLICATION NO.					DATE						
		2004				A1							 6174!				0030		
	CA	2507	455			AA		2004	0617	(CA 20	003-	2507	455		20	0031	121	
	WO	2004	0517	74		A2 20040617			WO 2003-US37336						20	0031	121		
	WO	2004	04051774 A3 2			2004	1125												
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
			CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
			GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	ΚP,	KR,	ΚZ,	LC,	
			LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NI,	NO,	
			NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ТJ,	
			TM,	TN,	TR,	TT,	TZ,	UA,	ŪĠ,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	zw		
		RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	
			BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	
			ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	IT,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	
			TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	TG
	EР	1565	957			A2		2005	0824		EP 20	003-	8124	43		2(0031:	121	
•		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,	
			IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	SK	-	
PRIOR	ITY	APP	LN.	INFO	. :					1	US 20	002-	4298	29P		P 20	0021	127	
						1	US 20	003-	4860	76P]	P 20030710							
							US 2003-617452				i	A 20030711							
							WO 2003-US37336				1	W 20031121							

OTHER SOURCE(S): MARPAT 141:9611

ED Entered STN: 28 May 2004

AB Disclosed are bioanodes comprising a quaternary ammonium treated Nafion polymer membrane and a dehydrogenase incorporated within the treated Nafion polymer. The dehydrogenase catalyzes the oxidation of an organic fuel and reduces an adenine dinucleotide. The ion conducting polymer membrane lies juxtaposed to a polymethylene green redox polymer membrane, which serves to electro-oxidize the reduced adenine dinucleotide. The bioanode is used in a fuel cell to produce high power densities.

IC ICM H01M004-90

ICS H01M004-96; H01M008-10; C12N011-08

INCL 429043000; 429044000; 429042000; 429030000; 429013000; 435180000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
Section cross-reference(s): 7, 38

```
IT
    Enzymes, uses
    RL: TEM (Technical or engineered material use); USES (Uses)
        (enzyme immobilization for use in biofuel cells and sensors)
    7439-89-6, Iron, uses 7439-97-6, Mercury, uses 7440-02-0, Nickel, uses
TΤ
     7440-06-4, Platinum, uses 7440-22-4, Silver, uses
                                                         7440-33-7, Tungsten,
    uses 7440-50-8, Copper, uses 7440-57-5, Gold, uses
     7782-42-5, Graphite, uses 11129-18-3, Cerium oxide
                                                          12597-68-1,
     Stainless steel, uses 12612-50-9, Molybdenum sulfide
     RL: MOA (Modifier or additive use); USES (Uses)
        (electron conductor; enzyme immobilization for use in biofuel cells and
       sensors)
     61-73-4, Methylene blue 92-31-9, Toluidine blue o 92-82-0D, Phenazine,
IT
             92-84-2, Phenothiazine 98-86-2, Acetophenone, uses 135-67-1,
     Phenoxazine 139-85-5, 3,4-Dihydroxybenzaldehyde 521-31-3, Luminol
     531-53-3, Azure A 531-55-5, Azure B 553-24-2, Neutral red 2381-85-3,
     Nile blue 2679-01-8, Methylene green 3625-57-8, Nile blue A
     7440-04-2D, Osmium, phenanthrolinedione 9003-01-4, Polyacrylic
           25013-01-8, Polypyridine 25233-30-1, Polyaniline 25233-34-5,
     Polythiophene 25265-76-3, Diaminobenzene 27318-90-7,
     1,10-Phenanthroline-5,6-dione 30604-81-0, Polypyrrole 37251-80-2,
     Toluidine blue 38096-29-6, Diaminopyridine 51878-01-4 54258-43-4,
     1,10-Phenanthroline-5,6-diol 68455-94-7D, Nitrofluorenone, derivs.
     74485-93-1, Poly(difluoroacetylene) 86090-24-6, Brilliant cresyl blue
     87257-37-2, Polythionine 103737-36-6, Toluene blue 104934-50-1,
     Poly(3-hexylthiophene) 126213-51-2, Poly(3,4-ethylenedioxythiophene)
     142189-51-3, Poly(thieno[3,4-b]thiophene 150645-85-5, Poly(neutral red)
     150645-86-6, Poly(methylene blue) 153312-51-7, Poly(3-(4-
     fluorophenyl)thiophene 161201-31-6 193265-88-2, Phenothiazin-5-ium,
     3-(dimethylamino)-7-(methylamino)-, chloride homopolymer 259737-85-4,
     Poly(3,4-ethylenedioxypyrrole) 308284-47-1, Benzo[a]phenoxazin-7-ium,
     5-amino-9-(diethylamino)-, sulfate (2:1) homopolymer 692776-93-5
     RL: CAT (Catalyst use); USES (Uses)
        (enzyme immobilization for use in biofuel cells and sensors)
     7631-86-9, Silica, uses
IT
     RL: MOA (Modifier or additive use); USES (Uses)
        (meso-porous, carbon-modified; enzyme
        immobilization for use in biofuel cells and sensors)
L62 ANSWER 10 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                       2004:119888 CAPLUS
ACCESSION NUMBER:
                        140:176638
DOCUMENT NUMBER:
TITLE:
                        Detoxification of nerve agents by surface
                        modified enzymes stabilized by
                        non-covalent immobilization on IDA silica
                        Singh, Alok; Pazirandeh, Mehran; Schoen, Paul E.;
INVENTOR(S):
                        Markowitz, Michael A.; Mauro, J. Matthew
                        The United States of America as Represented by The
PATENT ASSIGNEE(S):
                        Secretary of America, USA
                        U.S. Pat. Appl. Publ., 5 pp.
SOURCE:
                        CODEN: USXXCO
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO.
                                                                 DATE
                        _ _ _ _
                               -----
                                           -----
     US 2004029243
                         A1
                               20040212
                                          US 2000-725309
                                                                 20001129
     US 6869784
                         B2
                               20050322
     US 2003211586
                        A1
                               20031113
                                          US 2003-436602
                                                                 20030508
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PRIORITY APPLN. INFO.: US 2000-725309 A3 20001129 Entered STN: 13 Feb 2004 The present invention provides use of silica surface-immobilized AB thioesterase for detoxification of nerve agents. Enzymes, such as thioesterase, are modified by incorporating anchor sites for linking the enzymes to a target surface without destroying the catalytic activity of the enzymes. A a stable carrier to accommodate and bind the selected enzyme is constructed, and the enzyme is non-covalently linked to the carrier, generally through metal salts of iminodiacetate. The enzymes are genetically engineered to include stabilizing group selected from histidine and polyhistidine. The stabilized enzyme is attached to salt groups selected from the group consisting of metal salts of iminodiacetic acid, metal salts of nitrilotriacetic acid, and mixts. thereof on the surface of a particular inorg. carrier. ICM C12N009-00 IC ICS C12N011-08 INCL 435183000 4-3 (Toxicology) CC Section cross-reference(s): 7 TΤ Detoxification (biol.; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) Oxides (inorganic), uses IT RL: NUU (Other use, unclassified); USES (Uses) (ceramic particle as carrier for enzyme; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) ITAmphiphiles (copolymn. with amphiphiles containing salts to form vesicles; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) Decontamination IT Immobilization, molecular or cellular (detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) IT Enzymes, uses RL: NUU (Other use, unclassified); USES (Uses) (detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) ITProtein engineering (enzymes; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) ITSalts, uses RL: NUU (Other use, unclassified); USES (Uses) (for linking enzymes to silica surface; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) IT Chemical warfare agents (nerve gases; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica)

(of thioesterase; detoxification of nerve agents by surface

modified enzymes stabilized by non-covalent

Mutagenesis

IT

immobilization on IDA silica) 1304-28-5, Baria, uses 1310-53-8, Germania, uses 1344-28-1, Alumina, IT 7631-86-9, **Silica**, uses 13463-67-7, Titania, uses RL: NUU (Other use, unclassified); USES (Uses) (ceramic particle as carrier for enzyme; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) 71-00-1, Histidine, uses 26062-48-6, Polyhistidine IT RL: NUU (Other use, unclassified); USES (Uses) (enzyme stabilizing group comprising; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) ΙT 139-13-9D, Nitrilotriacetic acid, metal salts 142-73-4D, Iminodiacetic acid, alkoxysilane derivs., Cu2+ salts 7440-02-0D, Nickel, iminodiacetic or nitrilotriacetic acid salt 7440-48-4D, Cobalt, iminodiacetic or nitrilotriacetic acid salt 7440-50-8D, Copper, iminodiacetic or nitrilotriacetic acid salt 7440-66-6D, Zinc, iminodiacetic or nitrilotriacetic acid salt RL: NUU (Other use, unclassified); USES (Uses) (for linking enzymes to silica surface; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) TT 58943-36-5, Thioesterase RL: NUU (Other use, unclassified); USES (Uses) (from Escherichia coli; detoxification of nerve agents by surface modified enzymes stabilized by non-covalent immobilization on IDA silica) REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L62 ANSWER 11 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN 2003:509015 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 139:188235 TITLE: Direct Patterning of NiB on Glass Substrates Using Microcontact Printing and Electroless Deposition Geissler, Matthias; Kind, Hannes; Schmidt-Winkel, AUTHOR (S): Patrick; Michel, Bruno; Delamarche, Emmanuel CORPORATE SOURCE: Zurich Research Laboratory, IBM Research, Rueschlikon, 8803, Switz. Langmuir (2003), 19(15), 6283-6296 SOURCE: CODEN: LANGD5; ISSN: 0743-7463 PUBLISHER: American Chemical Society DOCUMENT TYPE: Journal English LANGUAGE: Entered STN: 04 Jul 2003 The authors investigate direct electroless deposition (ELD) of NiB on AB glass by microcontact-printing a reagent from an elastomeric stamp onto a glass substrate. There are three variants of this method depending on the reagent to be printed. ELD of a metal on a glass substrate necessitates the pretreatment of the glass with organic linkers that can bind a catalyst from solution The authors use colloidal Pd/Sn as catalyst and immobilize these particles via an amino-functionalized silane such as 3-(2-aminoethylamino)propyltrimethoxysilane (EDA-Si) grafted to the glass substrate. The first variant includes microcontact-printing EDA-Si onto glass to bind the colloidal Pd/Sn catalyst at well-defined locations on the substrate. Here, the stamp is first hydrophilized with an O2-based

plasma and then inked using different methods that include wet-inking, inking of the silane through the vapor phase, and contact inking. ELD of NiB initiates in those regions of the substrate that were previously

microcontact-printed. This approach entails the problem of inking and printing of an excess of silane, which can be washed away from the printed regions and can thereby induce the ELD of NiB grains adjacent to the desired pattern. In the second approach, the entire glass is uniformly derivatized with EDA-Si, and colloidal Pd/Sn particles are inked onto a stamp and microcontact-printed to activate the substrate where desired. These colloids do not diffuse on the substrate during printing and subsequent steps, allowing the formation of NiB patterns having excellent contrast and accuracy even over areas as large as 4'' in diameter Similar to the first method, inking and reusing stamps is inconvenient because the colloids are suspended in a solution of concentrated HCl and the stamp needs

to be

hydrophilized and covered with a thin layer of a polyelectrolyte to achieve homogeneous inking of the stamp with these colloids. The third and most promising approach is to derivatize the glass substrate homogeneously with both EDA-Si and Pd/Sn colloids and then deactivate the catalyst selectively prior to the ELD step by microcontact-printing eicosanethiol (ECT). In this case, it is possible to employ hydrophobic PDMS stamps, to reuse them, to optimize the inking and printing conditions, and to form high-quality NiB structures with lateral dimensions ranging from several hundred to below 1 μ m. This work suggests that microcontact printing (μ CP) and ELD can be combined in a variety of manners to provide interesting alternatives to conventional microfabrication methods that otherwise include the deposition of metals in a vacuum process and patterning methods based on photolithog.

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

IT Aluminoborosilicate glasses

RL: PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process)

(alkaline earth aluminoborosilicate, substrate; patterning of NiB by microcontact printing of linker or catalyst or deactivator on glass substrate followed by electroless deposition)

IT Silicone rubber, processes

RL: DEV (Device component use); PEP (Physical, engineering or chemical process); PYP (Physical process); PROC (Process); USES (Uses)

(di-Me, Sylgard 184, stamp; patterning of NiB by microcontact printing of linker or catalyst or deactivator on glass substrate followed by electroless deposition)

IT Coating process

(electroless; patterning of NiB by microcontact printing of linker or catalyst or deactivator on glass substrate followed by electroless deposition)

IT Lithography

(microcontact; patterning of NiB by microcontact printing of linker or catalyst or deactivator on glass substrate followed by electroless deposition)

IT Glass substrates

(patterning of NiB by microcontact printing of linker or catalyst or deactivator on glass substrate followed by electroless deposition)

IT Polyelectrolytes

(stamp treatment; patterning of NiB by microcontact printing of catalyst on glass substrate modified with amino-functionalized silane followed by electroless deposition)

linker or catalyst or deactivator on glass substrate followed

```
by electroless deposition)
     13373-97-2, 1-Eicosanethiol
IT
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); PROC (Process)
        (deactivator; patterning of NiB by microcontact printing of linker or
        catalyst or deactivator on glass substrate followed by
        electroless deposition)
     919-30-2, (3-Aminopropyl)triethoxysilane 1067-25-0D,
IT
     Propyltrimethoxysilane, polyethyleneimine-modified with
                                                               3069-29-2.
     3(2-Aminoethylamino)propylmethyldimethoxysilane 3179-76-8,
     3-Aminopropylmethyldiethoxysilane 9002-98-6D,
     trimethoxysilylpropyl-modified
                                     13497-18-2, Bis-[(3-
     triethoxysilyl)propyl]amine 41051-80-3, (3-Diethylaminopropyl)trimethoxy
             74956-86-8, N, N-Bis-[(3-trimethoxysilyl)propyl]ethylenediamine
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
    process); PROC (Process)
        (linker; patterning of NiB by microcontact printing of linker or
        catalyst or deactivator on glass substrate followed by
        electroless deposition)
     1760-24-3, 3-(2-Aminoethylamino)propyltrimethoxysilane
ΤТ
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)
        (linker; patterning of NiB by microcontact printing of linker or
        catalyst or deactivator on glass substrate followed by
        electroless deposition)
     11099-25-5P
                 94117-24-5P
IT
     RL: PEP (Physical, engineering or chemical process); PNU (Preparation,
     unclassified); PYP (Physical process); PREP (Preparation); PROC (Process)
        (patterning of NiB by microcontact printing of linker or
        catalyst or deactivator on glass substrate followed by
        electroless deposition)
     69898-76-6, Niposit 468
IT
     RL: PEP (Physical, engineering or chemical process); PYP (Physical
     process); PROC (Process)
        (patterning of NiB by microcontact printing of linker or
        catalyst or deactivator on glass substrate followed by
        electroless deposition)
     61840-27-5, Cartaretin F4
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (stamp treatment; patterning of NiB by microcontact printing of
        catalyst on glass substrate modified with amino-functionalized
        silane followed by electroless deposition)
                               THERE ARE 95 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         95
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L62 ANSWER 12 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                         2002:978586 CAPLUS
ACCESSION NUMBER:
                         138:35773
DOCUMENT NUMBER:
                         Methods and compositions for directed microwave
TITLE:
                         chemistry for accelerating chemical reactions
                         Martin, Mark
INVENTOR(S):
PATENT ASSIGNEE(S):
                         USA
                         U.S. Pat. Appl. Publ., 30 pp., Cont.-in-part of U.S.
SOURCE:
                         Provisional Ser. No. 237,192.
                         CODEN: USXXCO
DOCUMENT TYPE:
                         Patent
                         English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
```

PATENT NO.	KIND	DATE	APPLICATION NO.	. DATE			
	- -			-			
US 2002197645	A1	20021226	US 2001-968517		20011002		
US 2003082633	A1	20030501	US 2002-234092		20020905		
US 2004209303	A1	20041021	US 2004-842512		20040511		
US 2005191708	A1	20050901	US 2005-105460		20050414		
PRIORITY APPLN. INFO.:			US 2000-237192P	P	20001003		
			US 2001-968517	A2	20011002		
			US 2002-234092	A2	20020905		

ED Entered STN: 29 Dec 2002

The present invention concerns a novel means by which specific chosen reactions can be accelerated through the use of a new type of artificial enzyme. The invention allows specific reactions to occur at an accelerated rate, even in the presence of other non-chosen mols., which may be very similar in structure to the chosen reactant. The reactions may be stoichiometric or catalytic. Microwave energy is used to specifically accelerate chosen chemical reactions. The microwaves are directed to lossy materials that contain specific binding sites for the desired reactant (e.g., antigen, oligonucleotide, receptor ligand, enzyme substrate). Barium titanate, activated charcoal, and Bentonite clay gave microwave-targeted accelerated chemiluminescent reactions.

IC ICM G01N033-53 ICS C08K003-00

INCL 435007100; 204157680

CC 9-16 (Biochemical Methods)

Section cross-reference(s): 7

IT Antibodies and Immunoglobulins

Enzymes, uses

Zeolites (synthetic), uses

RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses)

(reactant-binding, composite with solid susceptible to dielec. heating; directed microwave chemical for accelerating (bio)chemical reactions)

IT Enzymes, uses

RL: CAT (Catalyst use); USES (Uses)

(synthetic; directed microwave chemical for accelerating (bio)chemical reactions)

IT 7439-89-6, Iron, uses 7440-03-1, Niobium, uses 7440-50-8,
 Copper, uses 12070-06-3, Tantalum carbide (TaC) 12136-78-6, Molybdenum
 silicide (MoSi2)

RL: CAT (Catalyst use); TEM (Technical or engineered material use); USES (Uses)

(mixture with insulator, susceptible to dielec. heating and containing specific binding agent for reactant; directed microwave chemical for accelerating (bio)chemical reactions)

IT 9003-01-4D, Acrylic acid polymer, compds. 9003-05-8,
 Polyacrylamide 9003-70-7, Poly(styrene-divinyl benzene) 9004-54-0D,
 Dextran, crosslinked, uses 9012-36-6, Agarose
 RL: TEM (Technical or engineered material use); USES (Uses)

(thermally-insulating coating; directed microwave chemical for accelerating (bio)chemical reactions)

L62 ANSWER 13 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2002:793841 CAPLUS

DOCUMENT NUMBER: 137:307013

TITLE: Non-enzymatic liposome-linked closely spaced array

electrodes assay (NEL-ELA) for detecting and

quantifying nucleic acids

INVENTOR(S): Bredehorst, Reinhard; Hintsche, Rainer; Heuberger,

Anton

PATENT ASSIGNEE(S): Fraunhofer-Gesellschaft zur Foerderung der Angewandten

Forschung e.V., Germany PCT Int. Appl., 124 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

SOURCE:

PATENT NO.					KIN	D	DATE		I	APP:	LICAT	ION :	NO.		D	ATE	
						-			-						-		
WO	2002	0817	39		A2		2002	1017	V	0	2002-	EP38	92		2	0020	408
WO	2002	0817	39		A3		2004	0129									
	W:	JP,	US														
	RW:	AT,	BE,	CH,	CY,	DE,	DK,	ES,	FI,	FR	, GB,	GR,	ΙE,	IT,	LU,	MC,	NL,
		PT,	SE,	TR													
EP	1409	728			A2		2004	0421	F	ΞP	2002-	7352	36		2	0020	408
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	, IT,	LI,	LU,	ΝL,	SE,	MC,	PT,
		ΙE,	FI,	CY,	TR												
PRIORITY	APP	LN.	INFO	. :					τ	JS	2001-	2821	64P		P 2	0010	409
									V	OV	2002-	EP38	92		W 2	0020	408

ED Entered STN: 18 Oct 2002

- The invention concerns target nucleic acids or amplicons thereof bound to immobilized capture oligonucleotides by mol. biol. reactions, are detected and quantified with affinity liposomes containing encapsulated electrochem. detectable reporter mols. susceptible to redox recycling and surface-attached affinity components capable of specifically binding to captured target nucleic acids or amplicons thereof in a structure restricted manner. Specifically bound affinity liposomes are lysed by temperature- or detergent-mediated mechanisms and released reporter mols. are quantitated via redox recycling using voltammetry in conjunction with a closely spaced array of thin film nobel metal electrodes. The quantity of released reporter mols. is a proportional measure of the quantity of target nucleic acids in the sample. For amplified assay procedures polymeric carrier mols. capable of binding multiple affinity liposomes or preformed complexes of affinity liposomes are utilized.
- IC ICM C12Q001-68
 - ICS G01N027-49
- CC 9-16 (Biochemical Methods)
 - Section cross-reference(s): 7
- IT Enzymes, uses
 - RL: ARG (Analytical reagent use); PRP (Properties); ANST (Analytical study); USES (Uses)
 - (inhibitors; non-enzymic liposome-linked closely spaced array electrodes assay (NEL-ELA) for detecting and quantifying nucleic acids)
- IT Enzymes, uses
 - Oligonucleotides
 - RL: ARG (Analytical reagent use); DEV (Device component use); ANST (Analytical study); USES (Uses)
 - (non-enzymic liposome-linked closely spaced array electrodes assay (NEL-ELA) for detecting and quantifying nucleic acids)
- IT 9002-89-5, Polyvinyl alcohol 9003-01-4, Polyacrylic acid
 - 9003-39-8, Polyvinylpyrrolidinone
 - RL: NUU (Other use, unclassified); USES (Uses)
 - (as carrier mols.; non-enzymic liposome-linked closely spaced array electrodes assay (NEL-ELA) for detecting and quantifying nucleic acids) 50-67-9, Serotonin, uses 50-76-0, Actinomycin D 51-61-6, Dopamine,
- uses 54-16-0, uses 66-97-7, Psoralen 120-12-7, Anthracene, uses 120-72-9, Indole, uses 120-80-9, Catechol, uses 123-30-8, p-Aminophenol 229-87-8, Phenanthridine 260-94-6, Acridine 288-32-4,

Imidazole, uses 495-99-8, Hydroxystilbamidine 7439-89-6, Iron, uses 7440-04-2, Osmium, uses 7440-18-8, Ruthenium, uses Chromium, uses 7440-50-8, Copper, uses 29249-00-1, Methoxytyramine RL: NUU (Other use, unclassified); USES (Uses) (non-enzymic liposome-linked closely spaced array electrodes assay (NEL-ELA) for detecting and quantifying nucleic acids) L62 ANSWER 14 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN 2002:578703 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 137:348377 The steady-state current at microcylinder electrodes TITLE: modified by enzymes immobilized in conducting or non-conducting material Somasundrum, M.; Aoki, K. AUTHOR (S): Faculty of Engineering, Department of Applied Physics, CORPORATE SOURCE: Fukui University, Fukui-shi, 910-8507, Japan SOURCE: Journal of Electroanalytical Chemistry (2002), 530(1-2), 40-46 CODEN: JECHES Elsevier Science B.V. PUBLISHER: DOCUMENT TYPE: Journal English LANGUAGE: Entered STN: 05 Aug 2002 A diffusion-kinetic model is presented for an enzyme-modified microcylinder electrode, where the enzyme reaction generates an electro-active product. Simple, approx. expressions are derived for the steady-state current in cases where the enzyme is immobilized in a metallically conducting matrix or a non-conducting matrix. The model is also extended to the case of a chemical sensor, which involves a conducting polymer without enzyme. The model is used to analyze steady-state signals for glucose produced by Pt-coated carbon fibers, on which glucose oxidase has been entrapped in poly(1,2-diaminobenzene). 7-7 (Enzymes) Electric current (biol.; diffusion-kinetic model addresses steady-state current at microcylinder electrodes modified by enzymes immobilized in conducting or non-conducting material) Carbon fibers, uses RL: NUU (Other use, unclassified); USES (Uses) (diffusion-kinetic model ests. kinetic parameters for glucose oxidase in poly(1,2-diaminobenzene) films on Pt-coated carbon fibers) Sensors (electrochem.; diffusion-kinetic model addresses steady-state current for conducting polymer-based chemical sensor) Microelectrodes (enzyme; diffusion-kinetic model addresses steady-state current at microcylinder electrodes modified by enzymes immobilized in conducting or non-conducting material) Immobilization, molecular or cellular (enzyme; diffusion-kinetic model ests. kinetic parameters for glucose oxidase in poly(1,2-diaminobenzene) films on Pt-coated carbon fibers) Enzymes, properties RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process) (immobilized; diffusion-kinetic model addresses steady-state current at microcylinder electrodes modified by enzymes immobilized in conducting or non-conducting material)

AB

CC

IT

IT

IT

IT

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IT

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TΤ
     Enzyme electrodes
        (microelectrodes; diffusion-kinetic model addresses steady-state
        current at microcylinder electrodes modified by
        enzymes immobilized in conducting or non-conducting
     7722-84-1, Hydrogen peroxide, processes
IT
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process): PROC (Process)
        (diffusion-kinetic model addresses H2O2-mediated current at
        microcylinder electrodes modified by enzymes
        immobilized in conducting or non-conducting material)
IT
     9001-37-0, Glucose oxidase
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PRP (Properties); PROC (Process)
        (diffusion-kinetic model ests. kinetic parameters for glucose oxidase
        in poly(1,2-diaminobenzene) films on Pt-coated carbon fibers)
     7440-06-4, Platinum, uses 25667-98-5, Poly(1,2-diaminobenzene)
IT
     RL: NUU (Other use, unclassified); USES (Uses)
        (diffusion-kinetic model ests. kinetic parameters for glucose oxidase
        in poly(1,2-diaminobenzene) films on Pt-coated carbon fibers)
                              THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                        40
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L62 ANSWER 15 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:
                        2002:51559 CAPLUS
DOCUMENT NUMBER:
                        136:119418
TITLE:
                        Modifying the surface of a substrate containing a
                        polymeric material, vaporizing surface modifying agent
                        onto a substrate, and apparatus
INVENTOR (S):
                        Filippou, Con; Gutowski, Wojciech Stanislaw; Proctor,
                        David; Spicer, Mark
PATENT ASSIGNEE(S):
                        Commonwealth Scientific and Industrial Research
                        Organisation, Australia
                        PCT Int. Appl., 73 pp.
SOURCE:
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
                        English
LANGUAGE:
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                        KIND
                               DATE
                                          APPLICATION NO.
                                                                  DATE
                               ------
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                                          -----
     WO 2002004552
                        A1 20020117 WO 2001-AU814
                                                                 20010706
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT,
            RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US,
            UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
    US 2002018860
                               20020214
                                           US 2001-780623
                         A1
                                                                  20010212
```

ED Entered STN: 18 Jan 2002

B2

A1

US 6706320

US 2004115354

PRIORITY APPLN. INFO.:

AB The modifying process comprises providing a solution of the surface modifying agent (e.g. crosslinkers, coupling agents, etc.) in a solvent or a surface

US 2003-700444

AU 2000-8590

US 2001-780623

20031105

A 20000706 A 20010212

20040316

20040617

modifying agent composition and optionally a carrier and subjecting the solution

to a zone of elevated temperature to vaporize the solvent or to vaporize the surface modifying agent composition and provide diffuse contact between the surface modifying agent and the surface of the substrate. Lupasol G 35 was used as a surface modifying agent in the treatment of automotive bumpers for enhanced adhesion to painting.

IC ICM C08J007-12

ICS C09J005-02; B05D003-00; B05D005-04

38-2 (Plastics Fabrication and Uses) CC

TΤ Burners

> (vaporizing chems.; for modifying surface of an automotive substrate containing a polymeric material)

1760-24-3, Z-6020 9002-98-6, Lupasol G 35 TΤ 26336-38-9,

30551-89-4, Poly(allylamine) Poly(vinylamine)

RL: TEM (Technical or engineered material use); USES (Uses)

(for modifying surface of an automotive substrate containing a polymeric material)

REFERENCE COUNT:

20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L62 ANSWER 16 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 2001:918926 CAPLUS

DOCUMENT NUMBER:

136:44556

TITLE:

SOURCE:

Microchip-based carbon dioxide gas sensor

INVENTOR (S):

Lee, Jae Seon; Shin, Jae Ho; Lee, Min Hyung; Nam,

Hakhyun; Cha, Geun Sig

PATENT ASSIGNEE(S):

I-Sens, Inc., S. Korea Eur. Pat. Appl., 20 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION: _____

EP 1164372 A2 20011219 EP 2001-113705	20010602
EP 1164372 A3 20040519	
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE	, MC, PT
IE, SI, LT, LV, FI, RO	
US 2002011408 A1 20020131 US 2001-877857	20010608
US 6746583 B2 20040608	
PRIORITY APPLN. INFO.: KR 2000-32050 A	20000612

Entered STN: 21 Dec 2001 ED

There is provided a microchip-based differential-type carbon dioxide gas AB sensor for detecting dissolved carbon dioxide levels. It functions with at least one working electrode composed of an unbuffered hydrogel membrane containing a certain amount of sodium bicarbonate and a pH-sensitive qas-permeable membrane; and a reference electrode composed of a buffered hydrogel membrane and a pH-sensitive gas-permeable membrane. unbuffered hydrogel membrane contains carbonic anhydrase, which reduces the time period for the hydration of carbon dioxide, thereby allowing the quick measurement of the level of carbon dioxide. In addition to being significantly improved in stabilization, sensing, and recovering time periods, the differential-type carbon dioxide gas sensor can be fabricated in small sizes and quickly measure levels of carbon dioxide dissolved in sample solution

IC G01N027-416

CC 72-3 (Electrochemistry) Section cross-reference(s): 6, 79

IT Hydration, chemical

(of carbon dioxide in microchip-based carbon dioxide gas sensor)

IT 9002-18-0, Agar 9002-89-5, Polyvinyl alcohol 9003-01-4, Polyacrylic acid 9003-39-8, Polyvinylpyrrolidone 9004-62-0, Hydroxyethyl cellulose 9004-65-3, Hydroxypropylmethyl cellulose 9011-14-7, Polymethylmethacrylate 79484-92-7, Methocel

RL: NUU (Other use, unclassified); USES (Uses)

(use as hygroscopic material in microchip-based carbon dioxide gas sensor with pH sensitive hydrogel membrane)

IT 1760-24-3, N-[3-(Trimethoxysilyl)propyl]ethylene diamine

RL: NUU (Other use, unclassified); USES (Uses)

(use as pH sensitive gas permeable membrane in microchip-based carbon dioxide gas sensor)

L62 ANSWER 17 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

2001:833647 CAPLUS

DOCUMENT NUMBER: TITLE:

135:355028

Method for the immobilization of recognition

substances using electrodeposition and application to

sensor preparation

INVENTOR(S):

Schuhmann, Wolfgang Asulab S.A., Switz.

PATENT ASSIGNEE(S): SOURCE:

PCT Int. Appl., 50 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent German

LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

								APPLICATION NO.									
	WO 2001086298					A1 20011115			WO 2001-CH279								
	W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	ΒZ,	CA,	CH,	CN,
		CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,
		HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	LS,
		LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	ΝZ,	PL,	PT,	RO,
		RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,
		VN,	YU,	ZA,	ZW,	AM,	AZ,	BY,	KG,	KZ,	MD,	RU,	TJ,	TM			
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZW,	ΑT,	BE,	CH,	CY,
		DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	TR,	BF,
		ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	NE,	SN,	TD,	TG		
DE	1002	2750			A1		2001	1122]	DE 2	000-	1002	2750		2	0000	510
CA	2408	470			AA		2002	1108	(CA 2	001-	24084	470		2	0010	508
EP	1282	819			A1		2003	0212		EP 2	001-	9252	66		2	0010	508
	R:										IT,	LI,	LU,	NL,	SE,	MC,	PT,
		ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	ΑL,	TR						
	2003															0010	508
US	2003	1530	61		A1		2003	0814	1	US 2	002-	2756	98		2	0021	108
PRIORIT	Y APP	LN.	INFO	. :					1	DE 2	000-	1002	2750	1	A 2	0000	510
									1	WO 2	001-	CH27	9	V	1 2	0010	508

ED Entered STN: 16 Nov 2001

AB The invention concerns cathodic or anodic electro-polymns. where sensor substances are deposited onto the metal surface along with the polymer layer, thus enzyme electrodes, arrays of sensor mols. are prepared Charged monomers are deposited also by changes of the pH. Polymers are carboxy-group containing homopolymers, or copolymers; recognition substances are enzymes, antibodies, antigens, haptens, oligonucleotides, peptides, proteins, lecithins, hormones, latex beads etc. Thus a glucose sensor was prepared by depositing onto a platinum electrode an electrodeposition resin

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and glucose oxidase.
    ICM G01N033-543
IC
     ICS C120001-68; B01J019-00; C120001-00; G01N033-545
     9-16 (Biochemical Methods)
    Section cross-reference(s): 3
IT
    Antibodies
    Antiqens
    DNA
      Enzymes, uses
    Haptens
    Hormones, animal, uses
    Lecithins
     Oligonucleotides
     Peptides, uses
     Proteins, general, uses
    Receptors
    RL: ARG (Analytical reagent use); DEV (Device component use); ANST
     (Analytical study); USES (Uses)
        (method for immobilization of recognition substances using
       electrodeposition and application to sensor preparation)
     79-10-7D, Acrylic acid, copolymers 1314-13-2, Zinc oxide, uses
TT
     7439-88-5, Iridium, uses 7439-97-6, Mercury, uses 7440-04-2, Osmium,
           7440-05-3, Palladium, uses 7440-06-4, Platinum, uses 7440-15-5,
     Rhenium, uses 7440-18-8, Ruthenium, uses 7440-22-4, Silver, uses
     7440-50-8, Copper, uses 7440-57-5, Gold, uses 7782-42-5,
     Graphite, uses 9003-01-4, Acrylic acid homopolymer 13463-67-7,
     Titanium dioxide, uses
     RL: DEV (Device component use); USES (Uses)
        (method for immobilization of recognition substances using
       electrodeposition and application to sensor preparation)
                               THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                        5
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L62 ANSWER 18 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                        2001:370076 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         135:195837
                        Synthesis and applications of bis(perfluorodecalin-1-
TITLE:
                         carbonyl) peroxide
                         Sawada, H.; Kurachi, M.; Takishita, K.; Tanedani, T.;
AUTHOR(S):
                         Terasawa, N.; Hayakawa, Y.; Kawase, T.
                         Department of Chemistry, Nara National College of
CORPORATE SOURCE:
                         Technology, Nara, Yamatokoriyama, Yata, 639-1080,
                         Japan
                         European Polymer Journal (2001), 37(7), 1409-1415
SOURCE:
                         CODEN: EUPJAG; ISSN: 0014-3057
                         Elsevier Science Ltd.
PUBLISHER:
DOCUMENT TYPE:
                         Journal
                        English
LANGUAGE:
     Entered STN: 23 May 2001
    New bis(perfluorodecalin-1-carbonyl) peroxide (PFDC-PO) was prepared by the
AB
     reaction of hydrogen peroxide with perfluorodecalin-1-carbonyl chloride,
     obtained from the chlorination of the corresponding carboxylic acid and
     phosphorus oxychloride in the presence of DMF. A kinetic study showed
     that PFDC-PO decomps. homolytically with a three-bond fission.
    Heptadecafluorodecahydro-1-naphthyl (F-DECALYL) end-capped oligomers were
    prepared by the reaction of PFDC-PO with monomers such as acrylic acid
     (ACA), N, N-dimethylacrylamide (DMAA), 3-methacryloxy-2-
     hydroxypropyltrimethylammonium chloride, and N-(1,1-dimethyl-3-
     oxobutyl)acrylamide (DOBAA) via a radical process. Interestingly, these
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F-DECALYL end-capped oligomers obtained were soluble in water and common organic solvents, although parent perfluorodecalin is insol. in water and common organic solvents except hexane. F-DECALYL end-capped ACA oligomer was easily soluble in water and reduced the surface tension of water effectively. Addnl., F-DECALYL end-capped ACA oligomer was found to enhance dramatically luminol chemiluminescence light yield compared to the corresponding linear fluoroalkylated ACA oligomer. F-DECALYL end-capped DMAA and DOBAA oligomers were insol. in water; however, these oligomers were soluble in common organic solvents. Especially, F-DECALYL end-capped DOBAA oligomer was found to form a monomol. film at the air-water interface. CC 35-3 (Chemistry of Synthetic High Polymers) polymn catalyst bis perfluorodecalin carbonyl peroxide prepn; stkinetics decompn catalyst bis perfluorodecalin carbonyl peroxide; luminol chemiluminescence enhancement perfluorodecalin end capped oligomer Decomposition kinetics TT Polymerization catalysts Solubility Surface pressure-area isotherms Surface tension (synthesis and applications of bis(perfluorodecalin-1-carbonyl) peroxide) IT 9003-01-4P, Acrylic acid homopolymer 25609-94-3P, 3-Methacryloxy-2-hydroxypropyltrimethylammonium chloride homopolymer 25897-89-6P, Poly[N-(1,1-Dimethyl-3-oxobutyl)acrylamide] 26793-34-0P, N, N-Dimethylacrylamide homopolymer RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (synthesis and applications of bis(perfluorodecalin-1-carbonyl) peroxide) REFERENCE COUNT: 36 THERE ARE 36 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L62 ANSWER 19 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2001:265492 CAPLUS DOCUMENT NUMBER: 134:304229 TITLE: Three dimensional array crosslinked films and methods of making such films INVENTOR (S): Raguse, Burkhard; Braach-Maksvytis, Vijoleta Lucija Bronislava PATENT ASSIGNEE(S): Commonwealth Scientific and Industrial Research Organisation, Australia SOURCE: PCT Int. Appl., 38 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: . PATENT NO. KIND APPLICATION NO. DATE DATE -------------------WO 2001025316 A1 20010412 WO 2000-AU1210 20001005 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,

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DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                20010412
     CA 2386241
                          AA
                                          CA 2000-2386241
                                                                    20001005
    EP 1244731
                          A1
                                20021002
                                           EP 2000-969074
                                                                    20001005
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
         R:
             IE, SI, LT, LV, FI, RO, MK, CY, AL
     JP 2003511251
                         Т2
                                20030325
                                            JP 2001-528479
                                                                    20001005
                                            AU 2000-78902
                                                                    20001005
    AU 777998
                                20041111
PRIORITY APPLN. INFO.:
                                            AU 1999-3264
                                                                 A 19991005
                                            WO 2000-AU1210
                                                                 W 20001005
    Entered STN: 13 Apr 2001
ED
     The nanoparticle film comprises a 3 dimensional crosslinked array of
AB
    nanoparticles and linker mols. The nanoparticle film is coherent, robust
     and self supporting. The film may be produced by adding linker mols. to a
     suspension of nanoparticles. The linker mols. form crosslinks between the
     nanoparticles. Prior to completion of the crosslinking reaction the
     crosslinked nanoparticles are separated from the suspension. Polyacrylic
     acid-crosslinked titania nanoparticle film (membrane supported) layer was
     deposited between 2 cystamine hydrochloride-crosslinked Au nanoparticle
     film layers to give a trilayer structure behaving as a capacitor.
     ICM C08J005-18
IC
     ICS C08F006-12; C30B001-02; H01L021-20
     76-10 (Electric Phenomena)
CC
     DNA
IT
       Enzymes, uses
     Fullerenes
     Polymers, uses
     Porphyrins
     Proteins, general, uses
     RL: TEM (Technical or engineered material use); USES (Uses)
        (linkers; self-supporting crosslinked nanoparticle films)
     56-17-7, Cystamine hydrochloride 107-15-3, 1,2-Ethylenediamine, uses
IT
     1191-62-4, 1,8-Octanedithiol 1892-29-1 1910-42-5, Methylviologen
                                  152996-47-9, 1,15-Pentadecanedithiol
     9003-01-4, Polyacrylic acid
     333971-30-5 333971-31-6
                                333971-32-7
     RL: TEM (Technical or engineered material use); USES (Uses)
        (linker; self-supporting crosslinked nanoparticle films)
     1303-00-0, Gallium arsenide (GaAs), uses 1306-23-6, Cadmium sulfide
IT
                   1306-24-7, Cadmium selenide (CdSe), uses
                                                               1306-25-8,
     (CdS), uses
     Cadmium telluride (CdTe), uses 1314-13-2, Zinc oxide (ZnO), uses
     1314-87-0, Lead sulfide (PbS) 1314-98-3, Zinc sulfide (ZnS), uses 1315-09-9, Zinc selenide (ZnSe) 7439-89-6, Iron, uses 7440-05-3
                                                              7440-05-3,
                     7440-06-4, Platinum, uses
                                                  7440-21-3, Silicon, uses
     Palladium, uses
     7440-22-4, Silver, uses 7440-48-4, Cobalt, uses 7440-50-8,
     Copper, uses 7440-57-5, Gold, uses 12781-95-2 13463-67-7, Titania,
            22398-80-7, Indium phosphide (InP), uses
                                                      50926-11-9, Indium tin
     RL: TEM (Technical or engineered material use); USES (Uses)
        (self-supporting crosslinked nanoparticle films)
IT
     9002-98-6
     RL: TEM (Technical or engineered material use); USES (Uses)
        (support; self-supporting crosslinked nanoparticle films)
                               THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         6
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L62 ANSWER 20 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                         2000:790686 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         133:331759
TITLE:
                         Method for detecting biological agents
                         Chen, Liaohai; Mcbranch, Duncan W.; Wang, Hsing-Lin;
INVENTOR (S):
```

Whitten, David G.

PATENT ASSIGNEE(S): The Regents of the University of California, USA

SOURCE: PCT Int. Appl., 38 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

```
DATE
    PATENT NO.
                       KIND
                                        APPLICATION NO.
    _____
                       ____
                              _____
                                          ______
    WO 2000066790
                        A1
                              20001109
                                         WO 2000-US12423
                                                                20000504
        W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
            CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
            IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA,
            MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI,
            SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY,
            KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
            DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
            CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                              20001109
                                       CA 2000-2340905
    CA 2340905
                        AA
                                                                20000504
                              20010509
                                        EP 2000-928892
    EP 1097242
                        A1
                                                                20000504
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
                              20030708
                                          US 2000-565589
                                                                20000504
                        B1
                                          AU 2000-47058
    AU 782254
                        B2
                              20050714
                                                                20000504
PRIORITY APPLN. INFO.:
                                          US 1999-132556P
                                                                19990505
                                                             P
                                          WO 2000-US12423
                                                             W 20000504
```

ED Entered STN: 10 Nov 2000

AB A sensor is provided including a polymer capable of having an alterable measurable property from the group of luminescence and elec. conductivity, the polymer having an intermediate combination of a recognition element, a tethering element and a property-altering element bound thereto and capable of altering the measurable property, the intermediate combination adapted for subsequent separation from the polymer upon exposure to an agent having an affinity for binding to the recognition element whereupon the separation of the intermediate combination from the polymer results in a detectable change in the alterable measurable property, and, a means of detecting said detectable change in the alterable measurable property.

IC ICM C12Q001-68
ICS C12Q001-70; G01N021-64; G01N033-00; G01N033-53; G01N033-531;

G01N033-533; G01N033-543; C07H021-02; C07H021-04; C12N015-00; B05D001-18; B01J013-00

CC 9-1 (Biochemical Methods)

IT Affinity

Bacteria (Eubacteria)

Biosensors

Cell

Dissolution

Electric conductivity

Energy transfer

Fluorescence

Fluorescent dyes

Fluorometry

Luminescence

Luminescence spectroscopy

Microorganism

Optical fibers

Polyelectrolytes

```
Sensors
     Separation
     Solutions
     Test kits
     Virus
        (method for detecting biol. agents)
     Antibodies
IT
     Antigens
     Coordination compounds
       Enzymes, uses
     Glycolipids
     Nucleic acids
     Oligonucleotides
     Peptide nucleic acids
     Plastics, uses
     Poly(arylenealkenylenes)
     Polyacetylenes, uses
     Polydiacetylenes
     Polymers, uses
     Polysaccharides, uses
     Proteins, general, uses
     Toxins
     RL: DEV (Device component use); USES (Uses)
        (method for detecting biol. agents)
     58-85-5, Biotin 71-00-1D, Histidine, copper complex
                                                               1910-42-5, Methyl
     viologen 7440-50-8D, Copper, histidine complex, uses
     9033-83-4, Polyphenylene 9055-67-8D, Poly(ADP-ribose) polymerase,
     DNA-binding domain 25067-54-3, Polyfuran 25067-54-3D, Polyfuran, derivs. 25067-58-7, Polyacetylene 25067-59-8, Polyvinyl carbazole
     25067-59-8D, Polyvinyl carbazole, derivs. 25233-30-1, Polyaniline
     25233-30-1D, Polyaniline, derivs. 25233-34-5, Polythiophene
     26009-24-5, Poly(p-phenylene vinylene) 30604-81-0, Polypyrrole
     30604-81-0D, Polypyrrole, derivs. 37758-47-7, Ganglioside GM1
     78675-98-6, Squaraine 96638-49-2, Poly(phenylene vinylene)
     96638-49-2D, Poly(phenylene vinylene), derivs. 103419-76-7,
     Poly(1,4-naphthalenediyl-1,2-ethenediyl) 125714-86-5 164658-06-4,
     Poly(2,5-pyridinediyl-1,2-ethenediyl) 189145-97-9, Poly(pyridinediyl-1,2-
     ethenediyl)
     RL: DEV (Device component use); USES (Uses)
        (method for detecting biol. agents)
                                THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                         9
                                RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L62 ANSWER 21 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN
                         2000:720661 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         134:53120
TITLE:
                         Adsorption of trypsin on silica-polymethylsiloxane and
                          its modified forms
                         Denisova, T. I.; Karpenko, G. F.; Khalyavka, T. A.;
AUTHOR (S):
                         Shvets, D. I.
CORPORATE SOURCE:
                         Inst. Sorbtsii i Problem Endoekologii, NAN Ukrainy,
                         Kiev, Ukraine
                         Ukrainskii Khimicheskii Zhurnal (Russian Edition)
SOURCE:
                          (2000), 66(1-2), 103-106
                         CODEN: UKZHAU; ISSN: 0041-6045
                         Institut Obshchei i Neorganicheskoi Khimii im. V. I.
PUBLISHER:
                         Vernadskogo NAN Ukrainy
DOCUMENT TYPE:
                         Journal
                         Russian
LANGUAGE:
    Entered STN: 13 Oct 2000
```

AΒ Adsorption-desorption of trypsin in aqueous salt solns. by organosilica sorbent and its Cu(II) -modified forms were examined It was demonstrated that modification in the studied conditions do not change porous structure parameters of the silica-polymethylsiloxane sorbent. It was noticed that Cu(II) present in composition of the sorbent increase adsorption and decelerate trypsin desorption into the aqueous salt solns. CC 7-7 (Enzymes) IT Immobilization, biochemical (enzyme; adsorption-desorption of trypsin on silica -polymethylsiloxane sorbent and Cu(II)-modified forms) 7440-50-8D, Copper, reaction products with silica-TT polymethylsiloxane, properties RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PRP (Properties); PROC (Process); USES (Uses) (adsorption-desorption of trypsin on silica-polymethylsiloxane sorbent and Cu(II) -modified forms) L62 ANSWER 22 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN 1999:727759 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 132:36099 TITLE: Preparation of perfluoroheptyl endcapped poly(acrylic acid) and surface tension of its aqueous solution Luo, Xiangdong; Li, Ruixia; Wu, Dacheng; Hsu, Shaw AUTHOR (S): Ling Textile College, Sichuan University, Chengdu, 610065, CORPORATE SOURCE: Peop. Rep. China Gaofenzi Xuebao (1999), (5), 623-626 SOURCE: CODEN: GAXUE9; ISSN: 1000-3304 PUBLISHER: Kexue Chubanshe DOCUMENT TYPE: Journal LANGUAGE: Chinese Entered STN: 17 Nov 1999 ED Five samples of perfluoroheptyl end-capped poly(acrylic acid) (FPAA) AB having a high surface activity were prepared by free radical polymerization of acrylic acid. The initiator was perfluorooctanoyl peroxide which produced the free radical of heptyl at 45°. The yields of FPAA were in the range of 10% to 25%. The weight-average mol. wts. were 2.62 x 104 to 7.99 x 104 and the polydispersity indexes were 1.97 to 5.64, depending on the concentration of the initiator in the monomer. The surface tension was about 15 mN/m at 30° for the aqueous solution of FPAA with the weight-average mol. weight 2.62 \times 104 in the concentration of about 0.01 g/mL. The mol. areas of FPAA at the air-water interface under the condition of critical micelle concentration were 3.2-4.6 nm2, which are obviously larger than the area per mol. of perfluorooctanoic acid (2.14 nm2) and much smaller than that of ordinary polyacrylic acid (83.4 nm2). The polymer mols. in the surface adsorbed layer form the brush structure because of the special effect of the perfluoroalkyl groups at the chain ends. 35-4 (Chemistry of Synthetic High Polymers) CC Molecular weight ITViscosity (preparation and properties of perfluoroheptyl end-capped poly(acrylic acid)) Polyelectrolytes

Surface tension

IT

(preparation of perfluoroheptyl end-capped poly(acrylic acid) and surface tension of its aqueous solution)

ITPolymerization

Polymerization catalysts

(radical; preparation of perfluoroheptyl end-capped poly(acrylic acid) and surface tension of its aqueous solution)

IT

RL: CAT (Catalyst use); USES (Uses)

(initiator; preparation of perfluoroheptyl end-capped

poly(acrylic acid) and surface tension of its aqueous solution)

9003-01-4DP, Poly(acrylic acid), perfluoroheptyl-terminated IT 34434-27-0DP, reaction products with poly(acrylic acid)

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation of perfluoroheptyl end-capped poly(acrylic acid) and surface tension of its aqueous solution)

L62 ANSWER 23 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1998:398403 CAPLUS

DOCUMENT NUMBER:

129:38129

TITLE:

Polyphenol-oxidizing enzymes-containing composition

and methods for treating porous article

INVENTOR(S):

Echigo, Takashi; Ohno, Ritsuko

PATENT ASSIGNEE(S):

Showa Denko K.K., Japan; Echigo, Takashi; Ohno,

Ritsuko

SOURCE:

PCT Int. Appl., 72 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA'	TENT NO.			KINI	DATE	APPLICATION NO.	DATE
WO	9824890 W: AU,			A1 CN,		1 WO 1997-JP3798 , NZ, RU, SG, US, VN	19971021
	RW: AT,	BE,	CH,	DE,	DK, ES, FI	, FR, GB, GR, IE, IT,	LU, MC, NL, PT, SE
JP	10218999			A2	1998081	3 JP 1997-142386	19970530
CA	2274248			AA	1998061	L CA 1997-2274248	19971021
AU	9745746			A1	1998062	9 AU 1997-45746	19971021 .
AU	736563			B2	2001080	2	
EP	953634			A1	1999110	B EP 1997-944193	19971021
	R: AT,	BE,	CH,	DE,	DK, ES, FR	, GB, GR, IT, LI, LU,	NL, SE, MC, PT,
	IE,	FI					
NZ	336135			Α	2000092	9 NZ 1997-336135	19971021
NO	9902742			Α	1999080	NO 1999-2742	19990604
US	20030175	65		A1	2003012	3 US 1999-319384	19990604
PRIORIT	Y APPLN.	INFO	.:			JP 1996-327252	A 19961206
						JP 1997-142386	A 19970530
						WO 1997-JP3798	W 19971021

ED Entered STN: 29 Jun 1998

AR Disclosed is a method for treating a porous article to increase its mol. weight by coating or impregnating with a composition containing an enzyme having a

polyphenol oxidation activity in an alkaline pH region, a phenolic compound, and/or

an aromatic amine compound The process is to improve its property such as strength, abrasion resistance, weather resistance, rust resistance, flame retardancy, antimicrobial property, preservative property, germicidal property, insecticidal property, insect repellency, antiviral property, biol. repellency, adhesion, sustained release of chems., coloring,

dimensional stability, cracking resistance, deodorant property, deoxygenation, humidification, water absorption, water repellency, surface smoothness, biol. affinity, ion exchange property, and formaldehyde absorption. The process also prevents chemical leaching and inorg. compds. from migrating onto the surface of the porous article. Polyphenol oxidase prepared from Myrothecium verrucaria was mixed with liquin sulfonate, cupric sulfate, ethylene diamine, and other compds. such as hinokitiol for the impregnation of cedar wood blocks at >pH 8 was demonstrated.

IC ICM C12N009-02

CC 7-8 (Enzymes)

Section cross-reference(s): 38, 43

Enzymes, uses TТ

RL: CAT (Catalyst use); NUU (Other use, unclassified); USES (Uses) (polyphenol-oxidizing; polyphenol-oxidizing enzymes-containing composition and

methods for treating porous article)

IT 57-06-7, Allyl isothiocyanate 91-20-3D, Naphthalene, derivs., uses 533-75-5, Tropolone 7429-90-5, Aluminum, uses 7439-95-4, Magnesium, uses 7439-98-7, Molybdenum, uses 7440-02-0, Nickel, uses 7440-22-4, Silver, uses 7440-38-2, Arsenic, uses 7440-42-8, Boron, uses 7440-47-3, Chrome, uses 7440-50-8, Copper, uses 7440-66-6, Zinc, uses 8062-15-5, Lignin sulfonate 9002-98-6 Lignin, uses 25322-68-3

RL: MOA (Modifier or additive use); USES (Uses)

(polyphenol-oxidizing enzymes-containing composition and methods for treating

porous article)

REFERENCE COUNT:

THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS 16 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L62 ANSWER 24 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1994:239633 CAPLUS

DOCUMENT NUMBER:

120:239633

TITLE:

Devices and methods for detection of an analyte based

upon light interference

INVENTOR (S):

Bogart, Gregory R.; Moddel, Garret R.; Maul, Diana M.; Etter, Jeffrey B.; Crosby, Mark; Miller, John B.;

Blessing, James; Kelley, Howard; Sandstrom, Torbjorn;

Stiblert, Lars

PATENT ASSIGNEE(S):

SOURCE:

Biostar, Inc., USA PCT Int. Appl., 208 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 14

PATENT INFORMATION:

PA.	FENT NO.			KINI)	DATE	}	APPLICATION NO.	DATE
					-				
WO	9403774			A1		1994	0217	WO 1993-US5673	19930610
	ra :W	, AU,	CA,	JP					
	RW: AT	, BE,	CH,	DE,	DK	, ES,	FR,	GB, GR, IE, IT, LU, MC,	NL, PT, SE
ΑU	9179004			A1		1992	1021	AU 1991-79004	19910320
ΑU	653940			B2		1994	1020		
ΕP	539383			A1		1993	0505	EP 1991-910056	19910320
ΕP	539383			В1		1996	0918		
	R: BE	, CH,	DE,	ES,	FR	, GB,	IT,	LI, LU, NL, SE	
JР	0550693	6		T2		1993	1007	JP 1991-509344	19910320
JР	3193373			B2		2001	0730	,	
ES	2094224			T3		1997	0116	ES 1991-910056	19910320

.тр	2001235473	2	A2		20010831	.TD	2000-287242		19910320
	9345360	,	A1		19940303		1993-45360		19930610
	07509565		T2		19951019		1994-505280		19930610
	3506703		B2		20040315	UF	1994-303200		1000010
					19960821		1002 015241		19930610
EP	727038	an an	A1	a -	19960821	EP	1993-915341		19930610
	R: ES, 1	FR, GB,	-	SE					
	1126278		A2		20010822	EP	2001-108521		19930610
EP	1126278		A 3		20011017				
	R: ES,	FR, GB,	IT,	SE					
JP	2002116208	3	A2		20020419	JP	2001-236186		19930610
JP	3507048		B2		20040315				
JP	200213949	3	A2		20020517	JP	2001-236144		19930610
JP	3456984		B2		20031014				
JP	200212260	1	A2		20020426	JP	2001-236166		20010803
JP	3673849		B2		20050720				
JР	2002122603	3	A2		20020426	JP	2001-236198		20010803
JР	3547723		B2		20040728				
	2005049350	5	A2		20050224	JР	2004-276389		20040924
	APPLN. II						1992-924343	Α	19920731
INTORITI							1991-910056	A	19910320
							1991-509344	A3	19910320
							1991-US1781	A	19910320
							1993-915341	A3	19930610
							1994-505280		19930610
							2001-236166		19930610
						WO	1993-US5673	W	19930610

ED Entered STN: 14 May 1994

Methods for analyzing an optical surface for an analyte of interest in a AB test sample and related instruments/devices are disclosed. The method entails the use of a thin-film optical immunoassay device whereby an analyte of interest is detected in a test sample through spectral changes in the light impinging on the surface prior to and after the binding of the analyte to a reactive substrate layer(s). The device includes a substrate which has a 1st color in response to light impinging thereon. The substrate also exhibits a 2nd color which is different from the 1st color. The 2nd color is exhibited in response to the same light when the analyte is present on the surface. Thus, SiO was vapor deposited on a polished monocryst. Si wafer to a thickness of 550 Å; the film had a golden interference color. The film was activated with N-(2-aminoethyl)-3-aminopropyltrimethoxysilane, coated with a DNP-albumin conjugate to a thickness of 40Å, rinsed, and dried. The coated wafer was used in a competitive immunoassay for DNP using goat anti-DNP antibody and an ellipsometer to measure the change in mass at the surface from the change in light intensity.

IC ICM G01B009-02

ICS G01N021-62

CC 9-1 (Biochemical Methods)

Section cross-reference(s): 79, 80

IT Ceramic materials and wares

Glass, oxide

Plastics

RL: ANST (Analytical study)

(attachment layer and optical thin film on substrate of, in interferometer for **chemical** anal.)

IT Silanes

RL: ANST (Analytical study)

(attachment layer containing, on interferometer for chemical anal.)

IT Reflectometers

(for chemical anal.)

IT Ellipsometers

```
(for chemical anal., attachment layers and optical thin films
        for)
     Semiconductor materials
IT
        (in interferometer, for chemical anal.)
TΤ
     Silazanes
     Silicates, uses
     Titanates
     RL: ANST (Analytical study)
        (optical thin film of, on ellipsometer for chemical anal.)
IT
    Latex
     Dendritic polymers
     Siloxanes and Silicones, uses
     RL: ANST (Analytical study)
        (optical thin film of, on interferometer for chemical anal.)
IT
     Electric conductors
        (oxide films, on interferometer for chemical anal.)
TΤ
     Polymers, uses
     RL: USES (Uses)
        (self-assembling, optical thin film of, on interferometer for
        chemical anal.)
IT
     Siloxanes and Silicones, uses .
     RL: USES (Uses)
        (aminoalkyl, branched, T-structured, films, on optical glass substrate
        in interferometer for chemical anal.)
TT
     Films
        (elec. conductive, oxide, on interferometer for chemical anal.)
     Optical materials
IT
        (films, on ellipsometer for chemical anal.)
IT
     Films
        (optical, on ellipsometer for chemical anal.)
IT
     Polymers, uses
     RL: USES (Uses)
        (star-branched, optical thin film of, on interferometer for
        chemical anal.)
     11105-01-4, Silicon oxynitride
IT
     RL: ANST (Analytical study)
        (antireflective film containing, on interferometer for chemical
        anal.)
IT 9002-98-6D, Polyethylenimine, trimethoxysilylpropyl-modified
     9003-17-2D, Polybutadiene, triethoxysilyl-modified 9003-53-6,
     Polystyrene 144856-48-4, TC 7A
     RL: ANST (Analytical study)
        (attachment layer of, on interferometer for immunoassay)
     1306-23-6, Cadmium sulfide, uses 7631-86-9, Silicon dioxide, uses
TT
     7782-40-3, Diamond, uses
     RL: USES (Uses)
        (composites with silicon, antireflective layer containing, on
        interferometer for chemical anal.)
     7440-47-3, Chromium, uses
IT
     RL: USES (Uses)
        (conductive layer of, on interferometer for chemical anal.)
IT
     7429-90-5, Aluminum, uses
     RL: USES (Uses)
        (film, on optical glass substrate in interferometer for chemical
        anal.)
IT
     7440-21-3, Silicon, uses
     RL: USES (Uses)
        (monocryst., attachment layer and optical thin film on substrate of, in
        interferometer for chemical anal.)
     7429-90-5D, Aluminum, alkoxides
                                       409-21-2, Silicon carbide, uses
IT
```

1314-23-4, Zirconium oxide, uses 12033-89-5, Silicon nitride, uses 13463-67-7, Titanium dioxide, uses RL: ANST (Analytical study) (optical thin film of, on ellipsometer for chemical anal.) 1760-24-3, N-(2-Aminoethyl)-3-aminopropyltrimethoxysilane IT RL: ANST (Analytical study) (silicon wafer activation with, for coating with antibody for immunoassav) L62 ANSWER 25 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN 1988:539199 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 109:139199 Adhesion promoting product containing alkoxysilane and TITLE: process for treating an integrated circuit substrate therewith Brewer, Terry; Flaim, Tony D.; Moss, Mary G. INVENTOR(S): PATENT ASSIGNEE(S): Brewer Science, Inc., USA SOURCE: Eur. Pat. Appl., 9 pp. CODEN: EPXXDW DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: APPLICATION NO. PATENT NO. KIND DATE DATE _____ --------------A2 EP 260976 EP 1987-308256 19880323 19870917 A3 19880803 EP 260976 R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE A 19900821 US 1986-908360 19860917 US 4950583 JP 63077123 JP 1987-223219 19870908 A2 19880407 US 1986-908360 A 19860917 PRIORITY APPLN. INFO.: Entered STN: 14 Oct 1988 A method of producing microelectronic components by photolithog. uses a AΒ new adhesion promoter from an alkoxysilane such as an aminophenylalkoxysilane, a nonamino aromatic alkoxysilane, a nonamino chloroalkylalkoxysilane or a nonamino acrylic alkoxysilane. Optionally, the adhesion promoter composition contains an adjuvant polymer and an adhesion promoting catalyst. The composition improves the adhesion of the microelectronic coating (e.g., from a photoresist) to the substrate. Thus, a composition containing methacryloxypropyltrimethoxysilane, isopropanol and ethanolamine produced improved adhesion of the pos. photoresist to a substrate e.g., Si and Si nitride. ICICM G03F007-02 74-6 (Radiation Chemistry, Photochemistry, and Photographic and Other CC Reprographic Processes) Section cross-reference(s): 76 780-69-8, Phenyltriethoxysilane 1646-59-9 **1760-24-3** IT 2530-83-8 4420-74-0, 3-Mercaptopropyltrimethoxysilane 7003-80-7, (p-Aminophenyl)triethoxysilane 13822-56-5, 3-Aminopropyltrimethoxysilane 27326-65-4 58068-97-6, 4,5-Dihydro-1-(3-(triethoxysilyl)propyl)-1H-70851-51-3, N-Trimethoxysilylpropylimidazole 74113-77-2 imidazole RL: USES (Uses) (adhesion promoting agent, for microelectronic component) 64-19-7, Acetic acid, uses and miscellaneous 141-43-5, Ethanolamine, IT uses and miscellaneous RL: USES (Uses)

(adhesion promoting catalyst, for microelectronic components) 2530-85-0 9002-98-6 9004-67-5, Methyl cellulose 54077-45-1

IT

RL: USES (Uses)

(adhesion promoting composition containing alkoxysilane and, for

microelectronic

components)

L62 ANSWER 26 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1988:506833 CAPLUS

DOCUMENT NUMBER:

109:106833

TITLE:

Use of modified titanium etc. species for the immobilization of bioactive compounds

enzymes

AUTHOR(S):

Kennedy, John F.; Cabral, Joaquim M. S.

CORPORATE SOURCE: SOURCE:

Dep. Chem., Univ. Birmingham, Birmingham, B15 2TT, UK Transition Metal Chemistry (Dordrecht, Netherlands)

(1987), 12(6), 481-93

CODEN: TMCHDN; ISSN: 0340-4285

DOCUMENT TYPE:

Journal; General Review

LANGUAGE:

English

Entered STN: 01 Oct 1988

A review with 26 refs., of enzyme immobilization on chemical modified AB transition metal oxides, especially Ti(IV) oxides. The topics discussed include

the crosslinking stabilization of enzymes immobilized on hydrous metal oxide type supports, enzyme immobilization on derivatized hydrous metal oxide type supports, and enzyme immobilization on organic polymers in association

with transition metal species.

7-0 (Enzymes) CC

Polymers, uses and miscellaneous IT

RL: USES (Uses)

(enzyme immobilization on transition metal oxides coated with)

IT Enzymes

RL: PROC (Process)

(immobilization of, on chemical modified transition metal oxides)

IT Immobilization, biochemical

(of enzymes, on chemical modified transition metal oxides)

13463-67-7DP, Titanium oxide (TiO2), chemical modified IT RL: SPN (Synthetic preparation); PREP (Preparation) (preparation and use in enzyme immobilization)

L62 ANSWER 27 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER:

1988:451291 CAPLUS

DOCUMENT NUMBER:

109:51291

TITLE:

Insoluble chemical composite material

suitable as a biocatalytic or affinity chromatographic

carrier and method for its preparation

INVENTOR(S):

Nakamura, Makoto; Kumazawa, Toshiaki; Kogure, Eriko

Olympus Optical Co., Ltd., Japan PATENT ASSIGNEE(S):

SOURCE:

Eur. Pat. Appl., 13 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 239079	A2	19870930	EP 1987-104341	19870324

EP 239079 A3 19880727

R: DE, FR, GB

JP 62226809 A2 19871005 JP 1986-69529 19860327 PRIORITY APPLN. INFO.: JP 1986-69529 A 19860327

ED Entered STN: 19 Aug 1988

- This material and method are described, where the material comprises a AB silicon dioxide-based carrier in the form of a particle, a fiber, or a tube, and polyethyleneimine, with an average mol. weight of 30,000 to 80,000, bonded to the carrier material by a silane coupling agent which is directly chemical bonded to the surface of the carrier material and comprises an epoxy-functional silane coupling agent to which the polyethyleneimine is directly bonded and is selected from the group consisting of β -(3,4-epoxycyclohexyl)ethyltrimethoxysilane, γ glycidoxypropyltrimethoxysilane, and γ-glycidoxypropylmethyldiethoxy silane. The silane coupling agent may not contain an epoxy group and the polyethyleneimine is covalently bonded to a linking agent, such as a dialdehyde compound like glutaraldehyde, covalently bonded with the silane coupling agent comprising an amino-functional silane coupling agent selected from the group consisting of N- β -(aminoethyl)- γ aminopropyltrimethoxysilane, N- β -(aminoethyl)- γ aminopropylmethyldimethoxysilane, \u03c4-aminopropyltriethoxysilane, and N-phenyl-γ-aminopropyltrimethoxysilane. A biochem., comprising an anti-human IgG monoclonal antibody or glucose oxidase, is bonded to an imino group and/or an amino group of the polyethyleneimine through a 2nd linking agent such as a dialdehyde like glutaraldehyde or such as a diepoxide compound like 1,4-butanediolglycidyl ether. This composite material has high moisture resistance and high alkali resistance. Bioreactor particles were prepared and filled into a column for a high-performance liquid chromatograph, and particles were prepared for an affinity-chromatog. carrier for human IgG separation
- IC ICM B01J020-32
- CC 9-3 (Biochemical Methods)

Section cross-reference(s): 80

IT Immunoglobulins

RL: ANST (Analytical study)

(G, separation of human, insol. **chemical** composite material as carriers for affinity chromatog. in)

IT Chromatography, column and liquid

(affinity, insol. chemical composite material as carriers for)

IT Reactors

(biocatalytic, insol. **chemical** composite material as carriers for)

IT Chromatography, column and liquid

(high-performance, insol. **chemical** composite material as carriers for)

IT 919-30-2D, γ -Aminopropyltriethoxysilane, reaction products with silica and polyethyleneimine 1760-24-3D, reaction products with silica and polyethyleneimine 2530-83-8D, γ -Glycidoxypropyltrimethoxysilane, reaction products with silica and polyethyleneimine 2897-60-1D, γ -Glycidoxypropylmethyldiethoxysilane, reaction products with silica and polyethyleneimine 3068-76-6D, reaction products with silica and polyethyleneimine 3069-29-2D, N- β -(Aminoethyl)- γ -aminopropylmethyldimethoxysilane, reaction products with silica and polyethyleneimine 3388-04-3D, β -(3,4-Epoxycyclohexyl)ethyltrimethoxysilane, reaction products with silica and polyethyleneimine

RL: ANST (Analytical study)

(coupling agent, insol. chemical composite material from, for biocatalytic or affinity chromatog. carriers)

IT 111-30-8D, Glutaraldehyde, reaction products with silica and

polyethyleneimine and silane coupling agent 1064-48-8 2425-79-8D, 1,4-Butanedioldiglycidyl ether, reaction products with silica and polyethyleneimine and silane coupling agent 9001-37-0D, Glucose oxidase, reaction products with silica and polyethyleneimine and silane coupling agent RL: ANST (Analytical study) (insol. chemical composite material from, for biocatalytic or affinity chromatog. carriers) 7631-86-9D, Silicon dioxide, reaction products with polyethyleneimine and silane coupling agent 7803-62-5D, Silane, derivs., reaction products with polyethyleneimine and silica 9002-98-6D, reaction products with silicon dioxide and silane coupling agent RL: ANST (Analytical study) (insol. composite material from, for biocatalytic or affinity chromatog. carriers) L62 ANSWER 28 OF 28 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 1981:605045 CAPLUS DOCUMENT NUMBER: 95:205045 Removal of heavy metals by chelate TITLE: resins Choi, Kyu Suck; Chung, Taek Sang; Chang, Byong Kwan; AUTHOR (S): Park, Young Ser CORPORATE SOURCE: Dep. Ind. Chem., Hanyang Univ., Seoul, S. Korea Polymer (Korea) (1981), 5(3), 194-202 SOURCE: CODEN: POLLDG; ISSN: 0379-153X DOCUMENT TYPE: Journal LANGUAGE: Korean Entered STN: 12 May 1984 Chelating resins for the absorption of heavy metals were prepared by treating silica gel or a cellulosic material, i.e., corncobs, with a coupling agent, polyethylenimine, and 2,4-TDI. In the case of resins on silica gel, the presence of silane couplers resulted in an increased adsorptivity for Pb2+ and Hg2+ but decreased adsorptivity for Cd2+. Resins prepared with methacrylate functional silanes showed the best adsorptivity for Hq2+. 37-3 (Plastics Fabrication and Uses) polyethylenimine TDI chelating resin; corncob carrier chelating resin; silica gel carrier chelating resin; silane coupler chelating resin; heavy metal chelating resin Corncob (in chelating resin manufacture, heavy metal adsorptivity in relation to) Coupling agents (silanes, in chelating resin preparation, heavy metal adsorptivity in relation to) Metals, reactions RL: RCT (Reactant); RACT (Reactant or reagent) (heavy, chelation of, carrier-supported resins for) 584-84-9D, reaction products with polyethylimine 9002-98-6D, reaction products with TDI RL: USES (Uses) (chelating resin manufacture from, for heavy metals) 1450-14-2 **1760-24-3** 2530-85-0 31681-13-7 RL: USES (Uses) (coupling agents, in chelate resin prepn, heavy metal adsorptivity in relation to)

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L63 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN ACCESSION NUMBER: 2004:394680 CAPLUS DOCUMENT NUMBER: 141:291503 Surface modification technology for bio-MEMS TITLE: AUTHOR (S): Shin, Dong-Sik; Kang, Chan-Ku; Kim, Jae-Kwon; Chung, Woo-Jae; Jang, Ki-Hoon; Lee, Yoon-Sik CORPORATE SOURCE: School of Chemical Engineering, Seoul National University, Seoul, 151-744, S. Korea Transducers '03, International Conference on SOURCE: Solid-State Sensors, Actuators and Microsystems, Digest of Technical Papers, 12th, Boston, MA, United States, June 8-12, 2003 (2003), Volume 2, 1746-1749. Institute of Electrical and Electronics Engineers: New York, N. Y. CODEN: 69FHV2; ISBN: 0-7803-7731-1 DOCUMENT TYPE: Conference LANGUAGE: English Entered STN: 16 May 2004 We have modified the surfaces of glass and stainless steel with polymers. AB The purposes of surface modification are for effective protein immobilization and increasing biocompatibility. The surface modified substrates were characterized by contact angle measurement, atomic force microscopy (AFM), XPS, and confocal fluorescence microscopy. Protein adsorption test was performed on the modified surfaces via enzyme-substrate reaction and fluorescence intensity measurement. We observed a drastic decrease of nonspecific adsorption of proteins when the surfaces are modified with hydrophilic polymers. CC 9-1 (Biochemical Methods) IT Reaction (Enzyme-substrate; surface modification technol. for bio-MEMS) REFERENCE COUNT: THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS 13 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L63 ANSWER 2 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN · 1997:343343 CAPLUS ACCESSION NUMBER: 127:78170 DOCUMENT NUMBER: TITLE: Efficient immobilization of proteins by modification of plate surface with polystyrene derivatives Suzuki, Noriko; Quesenberry, Michael S.; Wang, Judy AUTHOR (S): K.; Lee, Reiko T.; Kobayashi, Kazukiyo; Lee, Yuan Dep. Biology & McCollum-Pratt Inst., Johns Hopkins CORPORATE SOURCE: Univ., Baltimore, MD, 21218, USA Analytical Biochemistry (1997), 247(2), 412-416 SOURCE: CODEN: ANBCA2; ISSN: 0003-2697 PUBLISHER: Academic DOCUMENT TYPE: Journal LANGUAGE: English Entered STN: 31 May 1997 Immobilization of proteins on microplate wells by simple adsorption (e.g., AB for ELISA) is convenient, but it can be inefficient, especially if proteins are hydrophilic or small in size. This problem was alleviated by the use of polyvinylbenzyl lactonoylamide (PVLA). PVLA is strongly adsorbed to the hydrophobic well surface, and its lactonamide part can be oxidized with periodate to generate aldehydro groups. Proteins are then immobilized covalently to the aldehydo groups by reductive amination under mild

conditions. Using this method, henceforth termed the PVLA method, alkaline

phosphatase (AP) was immobilized to microplates six- to sevenfold greater than by simple adsorption (as measured by activity). Similarly, the activity of immobilized mannose-binding protein A (MBP-A) was 4- to 8-fold higher by the PVLA method than by simple adsorption. The PVLA-coated plates needed as little as 200 ng of MBP-A per well to have a sufficient amount of MBP-A immobilized for the measurement of binding of 125I-labeled mannosylated bovine serum albumin (125I-Man-BSA), but unmodified plates required as much as 20 $\mu g/well$ MBP-A to obtain the same response. Recommended conditions for the PVLA method are 40 μl of 2 mg/mL of PVLA for coating, 1 mM NaIO4 for the generation of the aldehydo groups, and a 2-h reductive amination at 37° between pH 8 and 9 for the protein ligation.

CC 9-16 (Biochemical Methods)

IT Immunoassay

(enzyme-linked immunosorbent assay; efficient immobilization of proteins by modification of plate surface with polystyrene derivs.)

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L63 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1995:924029 CAPLUS

DOCUMENT NUMBER: 124:28875

TITLE: Evaluation of the modifying influence of

arecanut on the garlic-modulated hepatic detoxication

system enzymes, sulfhydryl content, and

lipid peroxidation in mice

AUTHOR(S): Singh, Anjali; Rao, A. R.

CORPORATE SOURCE: School Life Sciences, Jawaharlal Nehru University, New

Delhi, India

SOURCE: Teratogenesis, Carcinogenesis, and Mutagenesis (1995),

15(3), 127-34

CODEN: TCMUD8; ISSN: 0270-3211

PUBLISHER: Wiley-Liss DOCUMENT TYPE: Journal LANGUAGE: English

ED Entered STN: 16 Nov 1995

AB This paper evaluates the potential effects of arecanut (Areca catechu, L.), an important ingredient of betel quid, on the garlic (Allium sativum, L.)-modulated activities of hepatic detoxication system enzymes, acid soluble sulfhydryl content, and lipid peroxidn. in mice. Mice were fed on either a normal diet or a diet containing 0.25%, 0.5%, or 1% (weight/weight) arecanut for

45 days. During the last 10 days of treatment garlic was orally administered at a dose level of 20 or 100 mg/kg/day. Significant modulation in the activities of phase I and phase II enzymes, -SH content, and malondialdehyde (MDA) level by garlic was observed Garlic-modulated alterations in glutathione S-transferase (GST) activity and -SH content were decreased, while cytochrome b5, cytochrome P 450, and MDA levels were further augmented by the arecanut plus garlic treatments.

CC 18-7 (Animal Nutrition)

L63 ANSWER 4 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1994:239520 CAPLUS

DOCUMENT NUMBER: 120:239520

TITLE: Preparation and characterization of ligand-modified

labeled liposomes for solid phase immunoassays

AUTHOR(S): Jones, Matthew A.; Singh, Anup; Kilpatrick,

Peter K.; Carbonell, Ruben G.

CORPORATE SOURCE: Dep. Chem. Eng., North Carolina State Univ., Raleigh,

NC, 27695-7905, USA

SOURCE: Journal of Liposome Research (1993), 3(3), 793-804

CODEN: JLREE7; ISSN: 0898-2104

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 14 May 1994

AB Small unilamellar vesicles conjugated with an enzyme label and with specific ligands for biol. mols. may prove to be useful as signal enhancement vehicles in the development of enzyme-linked immunoadsorbent assays and other detection applications. Bifunctional vesicles have been prepared by covalently attaching horseradish peroxidase (HRP) and monoclonal antibodies to the outside of the lipid bilayer. The reaction conditions were optimized to obtain 7-12 antibody mols. and 100-200 HRP mols. per vesicle. The enzyme retained 70-80% of its specific activity after immobilization with no apparent change in vesicle stability. These bifunctional vesicles were used in a noncompetitive immunoassay for D-Dimer, a fibrin dimer formed at the early stages of thrombogenesis. assay results using vesicles led to a detection limit for D-Dimer in human plasma which was five times lower than what was achieved using a conventional enzyme-antibody conjugate assay. HRP labeled (bifunctional) liposomes can also be used in competitive assays for the detection of small ligands in bulk solution HRP and biotin-conjugated vesicles were prepared and used in competitive assays for biotin in free solution The lowest detection limit for biotin using vesicles as the signal generation mechanism was a factor of 10 lower than what could be observed with a traditional biotin-HRP conjugate. A model has been developed for the competition between a small ligand in solution and a large ligand-conjugated vesicle for binding sites on a solid surface.

CC 9-10 (Biochemical Methods)
 Section cross-reference(s): 15

IT Immunoassay

(enzyme-linked immunosorbent assay, ligand-modified labeled liposomes for)

L63 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1979:536086 CAPLUS

DOCUMENT NUMBER: 91:136086

TITLE: Sequence-specific endonuclease Bgl I.

Modification of lysine and arginine residues

of the homogeneous enzyme

AUTHOR(S): Lee, Yan Hwa; Chirikjian, Jack G.

CORPORATE SOURCE: Med. Cent., Georgetown Univ., Washington, DC, 20007,

USA

SOURCE: Journal of Biological Chemistry (1979), 254(15),

6838-41

CODEN: JBCHA3; ISSN: 0021-9258

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 12 May 1984

AB The sequence-specific restriction endonuclease BglI from Bacillus globigii (RUB561) was purified to homogeneity as determined by denaturing polyacrylamide gel anal. The active form of the enzyme was a single polypeptide with a mol. weight of 32,000. The enzyme required Mg2+ in the reaction mixture and displayed a broad pH and monovalent cation requirement. BglI was not sensitive to SH-group reagents, but was affected by reagents that modify lysine and arginine residues. When lysine residues were modified by pyridoxal phosphate, both binding and catalysis were diminished whereas modification of arginine residues by 2,3-butanedione inhibited the enzyme activity but had no effect on its binding properties.

CC 7-2 (Enzymes)

L63 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN

ACCESSION NUMBER: 1978:559461 CAPLUS

DOCUMENT NUMBER: 89:159461

TITLE: Inactivation of phosphorylase b by potassium ferrate.

Identification of a tyrosine residue involved in the

binding of adenosine 5'-monophosphate

Lee, Young Moo; Benisek, William F.

CORPORATE SOURCE: Sch. Med., Univ. California, Davis, CA, USA

SOURCE: Journal of Biological Chemistry (1978), 253(15),

5460-3

CODEN: JBCHA3; ISSN: 0021-9258

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 12 May 1984

AB The site of reaction of K ferrate (K2FeO4) with rabbit muscle phosphorylase b was further characterized in an extension of previously

published studies reporting inactivation of the enzyme by this reagent. The tryptic peptide composed of residues 70-80 of the enzyme polypeptide chain contained a tyrosine residue which was chemical modified by ferrate and which was protected by AMP. The sequence of this peptide obtained from both untreated and ferrate-treated phosphorylase b was determined, and the results showed that tyrosine-75 was the residue with which ferrate reacts.

CC 7-5 (Enzymes) IT 13718-66-6

AUTHOR (S):

RL: BIOL (Biological study)

(phosphorylase b tyrosine modification by, enzyme

AMP binding site in relation to)

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